WEST Search History

Hide Items Restore Clear Cancel

DATE: Monday, October 18, 2004

Hide?	Set Name	Query	Hit Count
	DB=PGPB	,USPT,USOC,EPAB,JPAB,DWPI; PLUR=Y	ES; OP=ADJ
	L14	neuropeptide receptor	429
	L13	L12 AND neuropeptide receptor	78
	L12	530/300,350.CCLS.	17143
	L11	Rosen.IN.	6423
	L10	Rosen-C.IN.	19
	· L9	Rosen-C-A.IN.	590
	L8	Rosen-Craig.IN.	10
	L7	Rosen-Craig-A.IN.	657
	L6	Li.IN.	54090
	L5	Li-Y.IN.	4340
	L4	Li-Yi,IN.	240
	L3	Soppet.IN.	289
	L2	Soppet-D.IN.	5
	L1	(Soppet-Daniel.IN.)	2

END OF SEARCH HISTORY

Hit List

Clear Generate Collection Print Fwd Refs Bkwd Refs Generate OACS

Search Results - Record(s) 1 through 2 of 2 returned.

1. Document ID: US 20020081301 A1

Using default format because multiple data bases are involved.

L1: Entry 1 of 2

File: PGPB

Jun 27, 2002

PGPUB-DOCUMENT-NUMBER: 20020081301

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020081301 A1

TITLE: Cancer gene determination and therapeutic screening using signature gene sets

PUBLICATION-DATE: June 27, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Soppet, Daniel

Centreville

VA

US

US-CL-CURRENT: 424/155.1; 435/6, 514/1

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KMC | Draw Desc

2. Document ID: US 6338951 B1

L1: Entry 2 of 2

File: USPT

Jan 15, 2002

US-PAT-NO: 6338951

DOCUMENT-IDENTIFIER: US 6338951 B1

** See image for Certificate of Correction **

TITLE: G-protein parathyroid hormone receptor HLTDG74

DATE-ISSUED: January 15, 2002

INVENTOR-INFORMATION:

NAME CITY

STATE ZIP CODE

COUNTRY

Soppet; Daniel

Centreville

VA MD

Rosen; Craig A.

Gaithersburg Laytonsville

MD

Ruben; Steven M.

Olney

MD

US-CL-CURRENT: 435/69.1; 435/69.7, 514/12, 530/324, 530/350, 530/395, 530/402

ABSTRACT:

Li; Yi

Human G-protein parathyroid hormone (PTH) receptor polypeptides and DNA (RNA) encoding such polypeptides and a procedure for producing such polypeptides by

h eb b g ee ef e heh ef b

recombinant techniques is disclosed. Also disclosed are methods for utilizing such polypeptides for identifying antagonists and agonists to such polypeptides and methods of using the agonists and antagonists therapeutically to treat conditions related to the underexpression and overexpression of the PTH receptor receptor polypeptides. Also disclosed are diagnostic methods for detecting a mutation in the PTH receptor receptor nucleic acid sequences and detecting a level of the soluble form of the receptors in a sample derived from a host.

22 Claims, 10 Drawing figures Exemplary Claim Number: 1,16 Number of Drawing Sheets: 10

Full	Title Citation	Front	Review	Classification	Date	Reference					C	laims	KOMO	Drav
	***************************************	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	***************************************					**********	***************************************	**********	**********		*****	***************************************
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Clear	Gene	rate Co	llection	Print		Fwd Refs		Bkv	d Re	fs		Gene	arate	OACS
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Display Format: - Change Format

Previous Page Next Page Go to Doc#

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Hit List

Clear Generate Collection Print Fwd Refs Bkwd Refs Generate OACS

Search Results - Record(s) 1 through 5 of 5 returned.

1. Document ID: US 20040115625 A1, WO 200194629 A2, AU 200164559 A, US 20020081301 A1, US 20020102531 A1, US 20020102532 A1, US 20020110821 A1, US 20020115057 A1, US 20020115085 A1, US 20020150877 A1, US 20020165180 A1, US 20030165839 A1, EP 1358349 A2, JP 2004509612 W

Using default format because multiple data bases are involved.

L2: Entry 1 of 5

File: DWPI

Jun 17, 2004

DERWENT-ACC-NO: 2002-188264

DERWENT-WEEK: 200440

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TITLE: Screening for anti-neoplastic agent involves exposing cells to a chemical agent to be tested for anti-neoplastic activity, and determining a change in expression of a gene of a signature gene set

INVENTOR: AUGUSTUS, M; CARTER, K C; EBNER, R; ENDRESS, G; HORRIGAN, S; SOPPET, D R; WEAVER, Z; YOUNG, P E; SOPPET, D; YOUNG, P

PRIORITY-DATA: 2000US-245084P (November 1, 2000), 2000US-209473P (June 5, 2000), 2000US-209531P (June 5, 2000), 2000US-233133P (September 18, 2000), 2000US-233617P (September 18, 2000), 2000US-234009P (September 20, 2000), 2000US-234034P (September 20, 2000), 2000US-234052P (September 20, 2000), 2000US-234509P (September 22, 2000), 2000US-234567P (September 22, 2000), 2000US-234923P (September 25, 2000), 2000US-234924P (September 25, 2000), 2000US-235077P (September 25, 2000), 2000US-235082P (September 25, 2000), 2000US-235134P (September 25, 2000), 2000US-235280P (September 25, 2000), 2000US-235637P (September 26, 2000), 2000US-235638P (September 26, 2000), 2000US-235711P (September 27, 2000), 2000US-235720P (September 27, 2000), 2000US-235840P (September 27, 2000), 2000US-235863P (September 27, 2000), 2000US-236028P (September 28, 2000), 2000US-236032P (September 28, 2000), 2000US-236033P (September 28, 2000), 2000US-236034P (September 28, 2000), 2000US-236109P (September 28, 2000), 2000US-236111P (September 28, 2000), 2000US-236842P (September 29, 2000), 2000US-236891P (September 29, 2000), 2000US-237172P (October 2, 2000), 2000US-237173P (October 2, 2000), 2000US-237278P (October 2, 2000), 2000US-237294P (October 2, 2000), 2000US-237295P (October 2, 2000), 2000US-237316P (October 2, 2000), 2000US-237425F (October 3, 2000), 2000US-237598P (October 3, 2000), 2000US-237604P (October 3, 2000), 2000US-237606P (October 3, 2000), 2000US-237608P (October 3, 2000), 2000US-244867P (November 1, 2000), 2001US-0962436 (September 25, 2001), 2001US-0964824 (September 27, 2001), 2001US-0969708 (October 3, 2001), 2001US-0962832 (September 25, 2001), 2001US-0954456 (September 18, 2001), 2001US-0969347 (October 2, 2001), 2001US-0967768 (September 28, 2001), 2001US-0954531 (September 18, 2001), 2001US-0873367 (June 5, 2001), 2001US-0968007 (October 2, 2001)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
US 20040115625 A1	June 17, 2004		000	C12Q001/68
WO 200194629 A2	December 13, 2001	E	044	C12Q001/68
AU 200164559 A	December 17, 2001		000	
US 20020081301 A1	June 27, 2002		000	C12Q001/68
US 20020102531 A1	August 1, 2002		000	C12Q001/00
US 20020102532 A1	August 1, 2002		000	C12Q001/00

h e b b g e e e f e h eh ef b e

Jun 6, 2000

US 20020110821 A1	August 15, 2002		000	C12Q001/68
US 20020115057 A1	August 22, 2002		000	C12Q001/00
US 20020115085 A1	August 22, 2002		000	C12Q001/68
US 20020150877 A1	October 17, 2002		000	C12Q001/00
US 20020165180 A1	November 7, 2002		000	A61K038/17
US 20030165839 A1	September 4, 2003		000	C12Q001/68
EP 1358349 A2	November 5, 2003	E	000	C12Q001/68
JP 2004509612 W	April 2, 2004		083.	C12N015/09

A2 , JP 2004509612 W INT-CL (IPC): A61 K 31/00; A61 K 38/17; A61 K 39/395; A61 K 48/00; C07 H 21/04; C12 N 15/09; C12 Q 1/00; C12 Q 1/02; C12 Q 1/68; G01 N 33/15; G01 N 33/50; G01 N 33/566; G01 N 33/574

Full	Title	Citation Front	Review	Classification	Date	Reference		Claims	Kinic	Drain. Des
Г	2.	Document ID:	- US 60)71709 A			***************************************		*************	

File: DWPI

DERWENT-ACC-NO: 2000-411191

DERWENT-WEEK: 200035

L2: Entry 2 of 5

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TITLE: Detecting presence of neurotrophic factor or tyrosine kinase related oncogene receptor for diagnosing neurodegenerative diseases involves detecting tyrosine phosphorylation in a suspected sample

INVENTOR: KAPLAN, D; MARTIN-ZANCA, D; PARADA, L F; SOPPET, D

PRIORITY-DATA: 1992US-0890713 (May 29, 1992), 1991US-0668298 (March 14, 1991)

PATENT-FAMILY:

 PUB-NO
 PUB-DATE
 LANGUAGE
 PAGES
 MAIN-IPC

 US 6071709 A
 June 6, 2000
 027
 G01N033/567

INT-CL (IPC): $\underline{G01} \ \underline{N} \ \underline{33}/567$

ABSTRACTED-PUB-NO: US 6071709A

BASIÇ-ABSTRACT:

NOVELTY - Detecting a neurotrophic factor (NF) comprises contacting cells expressing tyrosine kinase(trk)-B-proto-oncogene receptor protein (TRP) with a putative NF and comparing the amount of phosphorylation of TRP in the cells, where an increase in phosphorylation compared with a control indicates presence of a NF.

DETAILED DESCRIPTION - An INDEPENDENT CLAIM is also included for a method of detecting a trk-proto-oncogene receptor protein in a sample comprising:

(1) contacting nerve growth factor (NGF) with a biological sample suspected of containing TRP and detecting any phosphorylation in the sample which indicates the presence of NGF.

USE - Identifying NF and TRP is useful for diagnosing degenerated neuronal diseases such as Alzheimer's and Huntington's disease in suspected patients, for diagnosing a tissue undergoing a neuronal regeneration and for designing compositions to treat neurodegenerative diseases.

Full | Title | Citation | Front | Review | Classification | Date | Reference | Classification | Claims | KWIC | Drawn Desc

3. Document ID: WO 9854963 A2, AU 9878120 A, EP 1039801 A1, JP 2002516573 W, US 20030092893 A1, EP 1428833 A2

L2: Entry 3 of 5

File: DWPI

Dec 10, 1998

DERWENT-ACC-NO: 1999-059865

DERWENT-WEEK: 200462

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TITLE: New isolated human genes and the secreted polypeptides they encode - useful for diagnosis and treatment of e.g. cancers, neurological disorders, immune diseases, inflammation or blood disorders

INVENTOR: BREWER, L A; CARTER, K; DILLON, P; EBNER, R; ENDRESS, G; FAN, P; FENG, P; FERRIE, A M; FISCHER, C; FLORENCE, C; FLORENCE, K; GREENE, J; HU, J; KYAW, H; LAFLEUR, D; LI, Y; MOORE, P A; NI, J; OLSEN, H; ROSEN, C; RUBEN, S; SHI, Y E; SOPPET, D; WEI, Y; YOUNG, P; YU, G; ZENG, Z; CARTER, K C; DILLON, P J; ENDRESS, G A; FISCHER, C L; GREENE, J M; LAFLEUR, D W; MORE, P A; OLSEN, H S; ROSEN, C A; RUBEN, S M; SHI, Y; SOPPET, D R

PRIORITY-DATA: 1997US-070923P (December 18, 1997), 1997US-048875P (June 6, 1997), 1997US-048876P (June 6, 1997), 1997US-048877P (June 6, 1997), 1997US-048878P (June 6, 1997), 1997US-048880P (June 6, 1997), 1997US-048881P (June 6, 1997), 1997US-048882P (June 6, 1997), 1997US-048883P (June 6, 1997), 1997US-048884P (June 6, 1997), 1997US-048885P (June 6, 1997), 1997US-048892P (June 6, 1997), 1997US-048893P (June 6, 1997), 1997US-048894P (June 6, 1997), 1997US-048895P (June 6, 1997), 1997US-048896P (June 6, 1997), 1997US-048897P (June 6, 1997), 1997US-048898P (June 6, 1997), 1997US-048899P (June 6, 1997), 1997US-048900P (June 6, 1997), 1997US-048901P (June 6, 1997), 1997US-048915P (June 6, 1997), 1997US-048916P (June 6, 1997), 1997US-048917P (June 6, 1997), 1997US-048949P (June 6, 1997), 1997US-048962P (June 6, 1997), 1997US-048963P (June 6, 1997), 1997US-048964P (June 6, 1997), 1997US-048970P (June 6, 1997), 1997US-048971P (June 6, 1997), 1997US-048972P (June 6, 1997), 1997US-048974P (June 6, 1997), 1997US-049019P (June 6, 1997), 1997US-049020P (June 6, 1997), 1997US-049373P (June 6, 1997), 1997US-049374P (June 6, 1997), 1997US-049375P (June 6, 1997), 1997US-057584P (September 5, 1997), 1997US-057627P (September 5, 1997), 1997US-057628P (September 5, 1997), 1997US-057629P (September 5, 1997), 1997US-057634P (September 5, 1997), 1997US-057635P (September 5, 1997), 1997US-057642P (September 5, 1997), 1997US-057643P (September 5, 1997), 1997US-057644P (September 5, 1997), 1997US-057645P (September 5, 1997), 1997US-057646P (September 5, 1997), 1997US-057647P (September 5, 1997), 1997US-057648P (September 5, 1997), 1997US-057649P (September 5, 1997), 1997US-057650P (September 5, 1997), 1997US-057651P (September 5, 1997), 1997US-057654P (September 5, 1997), 1997US-057661P (September 5, 1997), 1997US-057662P (September 5, 1997), 1997US-057666P (September 5, 1997), 1997US-057667P (September 5, 1997), 1997US-057668P (September 5, 1997), 1997US-057760P (September 5, 1997), 1997US-057761P (September 5, 1997), 1997US-057762P (September 5, 1997), 1997US-057763P (September 5, 1997), 1997US-057764P (September 5, 1997), 1997US-057765P (September 5, 1997), 1997US-057769P (September 5, 1997), 1997US-057770P (September 5, 1997), 1997US-057771P (September 5, 1997), 1997US-057774P (September 5, 1997), 1997US-057775P (September 5, 1997), 1997US-057776P (September 5, 1997), 1997US-057777P (September 5, 1997), 1997US-057778P (September 5, 1997), 1997US-049896P (June 6, 1997), 1998US-092921P (July 15, 1998), 1998US-094657P (July 30, 1998), 1998US-0205258 (December 4, 1998), 2001US-0023282 (December 20, 2001)

PATENT-FAMILY:

 PUB-NO
 PUB-DATE
 LANGUAGE
 PAGES
 MAIN-IPC

 WO 9854963 A2
 December 10, 1998
 E
 770
 A01N037/18

 AU 9878120 A
 December 21, 1998
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EP 1039801 A1	October 4, 2000	E	000	A01N037/18
JP 2002516573 W	June 4, 2002		914	C12N015/09
US 20030092893 A1	May 15, 2003		000	C07K016/00
EP 1428833 A2	June 16, 2004	E	000	C07K014/435

INT-CL (IPC): A01 N 37/18; A01 N 43/04; A61 K 31/711; A61 K 38/00; A61 K 38/17; A61 K 39/395; A61 K 48/00; A61 P 7/00; A61 P 25/00; A61 P 29/00; A61 P 35/00; A61 P 37/00; A61 P 43/00; C07 K 14/435; C07 K 14/47; C07 K 16/00; C07 K 16/18; C12 N 1/15; C12 N 1/19; C12 N 1/20; C12 N 1/21; C12 N 5/00; C12 N 5/06; C12 N 5/10; C12 N 15/00; C12 N 1/20; C12 N 15/00; C12 N 15/0

ABSTRACTED-PUB-NO: WO 9854963A BASIC-ABSTRACT:

An isolated nucleic acid molecule (NAM) (I) comprising a polynucleotide (PN) having a nucleotide sequence (NS) at least 95% identical to: (a) a PN fragment of one of a total of 207 defined human cDNA sequences given in the specification or a PN fragment of the cDNA sequence included in ATCC Deposit No. Z which is hybridisable to one of the 207 defined cDNA sequence; (b) a PN which is an (allelic) variant of one of the 207 defined cDNA sequences; (c) a PN encoding a biologically active polypeptide or a polypeptide fragment, domain or epitope of one of the 207 defined amino acid sequences given in the specification or a polypeptide fragment encoded by a cDNA sequence included in ATCC Deposit No. Z which is hybridisable to one of the defined cDNA sequences; (d) a PN which encodes a species homologue of one of the 207 defined polypeptides; or (e) a PN capable of hybridising under stringent conditions to any one of the PNs specified in (a)-(d), where the PN does not hybridise under stringent conditions to a sequence of only A residues or of only T residues. Also claimed are: (1) a recombinant vector comprising (I); (2) a method of making a recombinant host cell comprising (I); (3) a recombinant host cell produced by a method as in (2); (4) an isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from a polypeptide fragment (preferably having biological activity), domain, epitope, secreted form, full-length protein, (allelic) variant or species homologue of one of the 207 defined amino acid sequences or the encoded sequence included in ATCC Deposit No. Z; (5) an isolated antibody that binds specifically to an isolated polypeptide as in (4); (6) a recombinant host cell that expresses an isolated polypeptide as above; and (7) a gene corresponding to a cDNA sequence of the 207 defined amino acid sequences.

Note: From the disclosure 'ATCC Deposit No. Z' refers to the representative clones, each containing a subset of the defined cDNA sequences, which have been deposited with the ATCC. The deposit numbers are: ATCC 97979, 97974, 97975, 97976, 97977, 209007, 209008, 209009, 209010, 209011, 209080, 209081, 209082, 209083, 209084, 209085, 209511,.

USE The PNs and their corresponding secreted polypeptides are useful for preventing, treating or ameliorating medical conditions (claimed), e.g. by protein or gene therapy. Also pathological conditions can be diagnosed by determining the amount of the new polypeptides in a sample or by determining the presence of mutations in the new PNs (claimed). Specific uses are described for each of the 207 PNs, based on which tissues they are most highly expressed in, and include developing products for the diagnosis or treatment of cancer, tumours, neurodegenerative disorders, developmental abnormalities and foetal deficiencies, blood disorders, leukemias, diseases of the immune system, autoimmune diseases, hepatic and renal disease, lymphomas, inflammation, allergies, ischemic shock, Alzheimer's and cognitive disorders, schizophrenia, restenosis, prostate diseases, obesity, disorders involving osteoclasts such as osteoporosis, arthritis or malignancies, diseases of testes, lung or thymus, digestive/endocrine disorders, infections and AIDS. The polypeptides are also useful for identifying their binding partners (claimed).

Fuil	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KMAC	Draw, Desc

4. Document ID: JP 2003033192 A, WO 9639433 A1, AU 9526973 A, EP 832123 A1, JP 11507810 W, US 6030804 A, US 6338951 B1, US 20020086363 A1

L2: Entry 4 of 5

File: DWPI

Feb 4, 2003

DERWENT-ACC-NO: 1997-043068

DERWENT-WEEK: 200320

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TITLE: Human G-protein parathyroid hormone receptor, HLTDG74 - used to identify (ant) agonists, used in the treatment of hypo- or hyper-calcaemia, hypo- or hyper-phosphatemia, kidney stones, etc

INVENTOR: LI, Y; ROSEN, C A; RUBEN, S M; SOPPET, D R; SOPPET, D

PRIORITY-DATA: 1995WO-US07085 (June 5, 1995), 1995US-0468011 (June 6, 1995), 1999US-0236468 (January 25, 1999), 2001US-0996569 (November 30, 2001), 2002JP-0137833 (June 5, 1995)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
JP 2003033192 A	February 4, 2003		023	C12N015/09
WO 9639433 A1	December 12, 1996	E	062	C07K014/705
AU 9526973 A	December 24, 1996		000	C07K014/705
EP 832123 A1	April 1, 1998	E	000	C07K014/705
JP 11507810 W	July 13, 1999		056	C12N015/09
US 6030804 A	February 29, 2000		000	C12N015/12
US 6338951 B1	January 15, 2002	•	000	C07K014/72
US 20020086363 A1	July 4, 2002		000	C12P021/02

INT-CL (IPC): A61 K 35/76; A61 K 38/00; A61 K 39/395; A61 K 48/00; A61 P 3/14; A61 P 5/00; A61 P 13/12; A61 P 19/10; A61 P 43/00; C07 H 21/04; C07 K 14/705; C07 K 14/705; C07 K 14/72; C07 K 16/28; C12 N 1/21; C12 N 5/06; C12 N 5/10; C12 N 15/09; C12 N 15/12; C12 P 21/08; C12 Q 1/02; G01 N 33/15; G01 N 33/50; G01 N 33/567; C12 N 1/21; C12 R 1:19; C12 P 21/02; C12 P 21/0

ABSTRACTED-PUB-NO: US 6030804A

BASIC-ABSTRACT:

A novel isolated polynucleotide (I) comprises a member selected from: (a) a polynucleotide of 1914 bp encoding the polypeptide of 541 residues given in the specification; (b) a polynucleotide encoding a mature polypeptide encoding by the DNA deposited as ATCC 97186; (c) a polynucleotide capable of hybridising to, and which is at least 70% identical to the nucleotide sequence of (a) or (b); and (d) a polynucleotide fragment of the nucleotide sequences of (a), (b) or (c).

USE - The cpds. of (6) may be used for the treatment of patients which need to activate or inhibit a G-protein coupled receptor (claimed). Mutations in (I) or the corresp. protein may be identified by sequence analysis. Agonists cpds. may be used to prevent and/or treat hypocalcaemia, hyperphosphatemia, hypoparathyroidism and chronic tetany by stimulating an increase in serum calcium levels. Antagonist cpds. may be used to treat and/or prevent osteoporosis, hypercalcaemia, hypoparathyroidism, hypophosphatemia, kidney stone and nephrolithiasis.

ABSTRACTED-PUB-NO:

US 6338951B EQUIVALENT-ABSTRACTS:

A novel isolated polynucleotide (I) comprises a member selected from: (a) a

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Apr 1, 1993

Record List Display

polynucleotide of 1914 bp encoding the polypeptide of 541 residues given in the specification; (b) a polynucleotide encoding a mature polypeptide encoding by the DNA deposited as ATCC 97186; (c) a polynucleotide capable of hybridising to, and which is at least 70% identical to the nucleotide sequence of (a) or (b); and (d) a polynucleotide fragment of the nucleotide sequences of (a), (b) or (c).

USE - The cpds. of (6) may be used for the treatment of patients which need to activate or inhibit a G-protein coupled receptor (claimed). Mutations in (I) or the corresp. protein may be identified by sequence analysis. Agonists cpds. may be used to prevent and/or treat hypocalcaemia, hyperphosphatemia, hypoparathyroidism and chronic tetany by stimulating an increase in serum calcium levels. Antagonist cpds. may be used to treat and/or prevent osteoporosis, hypercalcaemia, hypoparathyroidism, hypophosphatemia, kidney stone and nephrolithiasis.

A novel isolated polynucleotide (I) comprises a member selected from: (a) a polynucleotide of 1914 bp encoding the polypeptide of 541 residues given in the specification; (b) a polynucleotide encoding a mature polypeptide encoding by the DNA deposited as ATCC 97186; (c) a polynucleotide capable of hybridising to, and which is at least 70% identical to the nucleotide sequence of (a) or (b); and (d) a polynucleotide fragment of the nucleotide sequences of (a), (b) or (c).

USE - The cpds. of (6) may be used for the treatment of patients which need to activate or inhibit a G-protein coupled receptor (claimed). Mutations in (I) or the corresp. protein may be identified by sequence analysis. Agonists cpds. may be used to prevent and/or treat hypocalcaemia, hyperphosphatemia, hypoparathyroidism and chronic tetany by stimulating an increase in serum calcium levels. Antagonist cpds. may be used to treat and/or prevent osteoporosis, hypercalcaemia, hypoparathyroidism, hypophosphatemia, kidney stone and nephrolithiasis.

US20020086363A

A novel isolated polynucleotide (I) comprises a member selected from: (a) a polynucleotide of 1914 bp encoding the polypeptide of 541 residues given in the specification; (b) a polynucleotide encoding a mature polypeptide encoding by the DNA deposited as ATCC 97186; (c) a polynucleotide capable of hybridising to, and which is at least 70% identical to the nucleotide sequence of (a) or (b); and (d) a polynucleotide fragment of the nucleotide sequences of (a), (b) or (c).

USE - The cpds. of (6) may be used for the treatment of patients which need to activate or inhibit a G-protein coupled receptor (claimed). Mutations in (I) or the corresp. protein may be identified by sequence analysis. Agonists cpds. may be used to prevent and/or treat hypocalcaemia, hyperphosphatemia, hypoparathyroidism and chronic tetany by stimulating an increase in serum calcium levels. Antagonist cpds. may be used to treat and/or prevent osteoporosis, hypercalcaemia, hypoparathyroidism, hypophosphatemia, kidney stone and nephrolithiasis.

WO 9639433A

Full | Title | Citation | Front | Review | Classification | Date | Reference | Claims | Claims | KWC | Draws Designation | Date | Reference | Claims | Claims | KWC | Draws Designation | Date | Reference | Claims | Claims | KWC | Draws Designation | Date | Reference | Claims | Claims | KWC | Draws Designation | Date | Reference | Claims | Claims | KWC | Draws Designation | Date | Reference | Claims | Claims | KWC | Draws Designation | Date | Reference | Claims | Claims | KWC | Draws Designation | Date | Reference | Claims | Claims | KWC | Draws Designation | Date | Reference | Claims | Claims | KWC | Draws Designation | Date | Reference | Claims | Claims | KWC | Draws Designation | Date | Reference | Claims | Claims | KWC | Draws Designation | Date | Reference | Claims | Claims | Claims | KWC | Draws Designation | Date | Reference | Claims | Claim

File: DWPI

DERWENT-ACC-NO: 1993-152039

DERWENT-WEEK: 200173

L2: Entry 5 of 5

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TITLE: Complexes of tyrosine receptor kinase - with nerve growth factors, used for study, diagnosis and treatment of neuro-degenerative diseases

h eb b g ee ef e h eh ef b

Record List Display

INVENTOR: KAPLAN, D; MARTIN-ZANCA, D; PARADA, L; SOPPET, D

PRIORITY-DATA: 1991US-0890713 (March 14, 1991)

PATENT-FAMILY:

 PUB-NO
 PUB-DATE
 LANGUAGE
 PAGES
 MAIN-IPC

 US N7890713 N
 April 1, 1993
 071
 C12N000/00

INT-CL (IPC): C12N 0/00

ABSTRACTED-PUB-NO: US 7890713A

BASIC-ABSTRACT:

The following are disclosed: (A) complex comprising nerve growth factor (NGF) and tyrosine receptor kinase (trk) proto-oncogene protein, where the complex is free of protein with which it is naturally associated; (B) a complex of neurotrophin-3 (NT-3) or brain-derived neurotrophic factor (BDNF) and trkB-proto-oncogene protein, where the complex is free of protein with which it is naturally associated; (C) a method of detecting the NGF:trk-proto-oncogene receptor complex, NT-3:trkB proto-oncogene receptor complex or BDNF:trkB proto-oncogene receptor complex in a sample which comprises contacting the sample with an antibody that binds specifically with NGF, NT-3, BDNF, trk- or trkB-proto-oncogene receptor protein of the complex, a positive immunological reaction indicating the presence of the complex; (D) a method of diagnosing degenerative neuronal diseases in a patient, which comprises contacting a sample of diseased tissue with an antibody that binds with one of the complexes as in (C) and detecting complex formation; (E) a method of diagnosing a tissue undergoing neuronal regeneration in a patient, which comprises contacting a sample of the tissue with an antibody that binds to one of the tissues as in (C) and assaying for the presence of resulting complex; (F) a method of detecting NGF, NT-3 or BDNF in a sample which comprises contacting the sample with trk or trkB-proto-oncogene receptor protein and detecting the presence of bound NGF, NT-3 or BDNF; (G) a method of detecting trk or trkB-proto-oncogene receptor protein in a sample using NGF, NT-3 or BDNF as a binding agent; (H) a method of detecting neurotrophic factor receptor/ligand complexes that are structurally and functionally related to trk and NGF comprising using the methods described above for detecting trk:NGF, trkB:NT-3 and trkB:BDNF complexes.

USE - The methods can be used for the diagnosis of neurodegenerative diseases that affect NGF-dependent neurons such as Alzheimer's and Huntington's disease. The methods can also be used to study nerve survival and regeneration and to develop therapeutic methods for treating such diseases

The following are disclosed: (A) complex comprising nerve growth factor (NGF) and tyrosine receptor kinase (trk) proto-oncogene protein, where the complex is free of protein with which it is naturally associated; (B) a complex of neurotrophin-3 (NT-3) or brain-derived neurotrophic factor (BDNF) and trkB-proto-oncogene protein, where the complex is free of protein with which it is naturally associated; (C) a method of detecting the NGF:trk-proto-oncogene receptor complex, NT-3:trkB proto-oncogene receptor complex or BDNF:trkB proto-oncogene receptor complex in a sample which comprises contacting the sample with an antibody that binds specifically with NGF, NT-3, BDNF, trk- or trkB-proto-oncogene receptor protein of the complex, a positive immunological reaction indicating the presence of the complex; (D) a method of diagnosing degenerative neuronal diseases in a patient, which comprises contacting a sample of diseased tissue with an antibody that binds with one of the complexes as in (C) and detecting complex formation; (E) a method of diagnosing a tissue undergoing neuronal regeneration in a patient, which comprises contacting a sample of the tissue with an antibody that binds to one of the tissues as in (C) and assaying for the presence of resulting complex; (F) a method of detecting NGF, NT-3 or BDNF in a sample which comprises contacting the sample with trk or trkB-proto-oncogene receptor protein and detecting the presence of bound NGF, NT-3 or BDNF; (G) a method of detecting trk or trkB-proto-oncogene receptor protein in a sample using NGF, NT-3 or BDNF as a binding agent; (H) a method of detecting neurotrophic factor

receptor/ligand complexes that are structurally and functionally related to trk and NGF comprising using the methods described above for detecting trk:NGF, trkB:NT-3 and trkB:BDNF complexes.

USE - The methods can be used for the diagnosis of neurodegenerative diseases that affect NGF-dependent neurons such as Alzheimer's and Huntington's disease. The methods can also be used to study nerve survival and regeneration and to develop therapeutic methods for treating such diseases ABSTRACTED-PUB-NO:

US N7890713N EQUIVALENT-ABSTRACTS:

Full Title Citation Front Review	Classification Date Reference		Claims KW	C Draw Des
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Terms	Docum	nents		

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Search Results - Record(s) 1 through 10 of 10 returned.

1. Document ID: US 20020146778 A1

Using default format because multiple data bases are involved.

L8: Entry 1 of 10

File: PGPB

Oct 10, 2002

RULE-47

PGPUB-DOCUMENT-NUMBER: 20020146778

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020146778 A1

TITLE: Pineal gland specific gene-1

PUBLICATION-DATE: October 10, 2002

INVENTOR-INFORMATION:

NAME CITY

Y STATE COUNTRY

He, Wei Wu

Rosen, Craig

Columbia MD US Laytonsville MD US

US-CL-CURRENT: 435/69.4; 435/320.1, 435/325, 530/399, 536/23.5

Full: Title Citation Front Review Classification Date Reference Sequences Attachments Claims KMC Draw Des

2. Document ID: US 20020086314 A1

L8: Entry 2 of 10

File: PGPB

Jul 4, 2002

PGPUB-DOCUMENT-NUMBER: 20020086314

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020086314 A1

TITLE: Colon specific genes and proteins

PUBLICATION-DATE: July 4, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Yu, Guo-Liang Berkeley CA US

Rosen, Craig Laytonsville MD US

 $\text{US-CL-CURRENT: } \underline{435/6}; \ \underline{435/196}, \ \underline{435/320.1}, \ \underline{435/325}, \ \underline{435/69.1}, \ \underline{435/7.23}, \ \underline{536/23.2}$

ABSTRACT:

Human colon specific gene polypeptides and DNA (RNA) encoding such polypeptides and a procedure for producing such polypeptides by recombinant techniques is disclosed. Also disclosed are methods for utilizing such polypucleotides or polypeptides as a

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diagnostic marker for colon cancer and as an agent to determine if colon cancer has metastasized. Also disclosed are antibodies specific to the colon specific gene polypeptides which may be used to target cancer cells and be used as part of a colon cancer vaccine. Methods of screening for agonists and antagonists for the polypeptide and therapeutic uses of the antagonists are disclosed.

Full * Title * Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KWC | Draw Des

3. Document ID: US 20020042119 A1

L8: Entry 3 of 10

File: PGPB

Apr 11, 2002

PGPUB-DOCUMENT-NUMBER: 20020042119

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020042119 A1

TITLE: Novel Metalloproteinases

PUBLICATION-DATE: April 11, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Ni, Jian	Germantown	MD	US	
Ruben, Steve	Olney	MD	US	
Brewer, Laurie	St. Paul	MD	US	
Gentz, Reiner	Rockville	MD	US	
Rosen, Craiq	Laytonsville	MD	US	

US-CL-CURRENT: <u>435/219</u>; <u>435/320.1</u>, <u>435/325</u>, <u>435/69.1</u>, <u>536/23.2</u>

ABSTRACT:

The present invention relates to novel metalloproteinase-like proteins. In particular, isolated nucleic acid molecules are provided encoding the human TACE-like and matrilysin-like proteins. TACE-like and matrilysin-like polypeptides are also provided as are vectors, host cells and recombinant methods for producing the same. The invention further relates to screening methods for identifying agonists and antagonists of TACE-like and matrilysin-like activity. Also provided are diagnostic methods for detecting cancer and therapeutic methods for cancer and other disorders characterized by an over or under production of these metalloprofeinases.

Full Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draw Desc

4. Document ID: US 6608182 B1

L8: Entry 4 of 10

File: USPT

Aug 19, 2003

US-PAT-NO: 6608182

DOCUMENT-IDENTIFIER: US 6608182 B1

TITLE: Human vascular endothelial growth factor 2

DATE-ISSUED: August 19, 2003

h e b b g e e e f e h eh ef b

COUNTRY

INVENTOR-INFORMATION:

....

NAME CITY STATE ZIP CODE

Rosen; Craiq Laytonsville MD Hu; Jing-Shan Gaithersburg MD

Cao; Liang Monmouth Terrace HK

US-CL-CURRENT: 530/399; 435/243, 435/320.1, 435/325, 435/69.1, 435/69.4, 530/300, 530/350, 530/402

ABSTRACT:

The present invention relates to polypeptides comprising amino acids 85 to 165 of SEQ ID NO:2, as well as polynucleotides which encode these polypeptides. Also provided are methods of treatment using these polypeptides.

21 Claims, 15 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 15

	Draw, Des	KMC	Claims			Reference	Date	Classification	Review	Front	Citation	Title	Full
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5. Document ID: US 6337195 B1													

1... J. Document ID. 08 033/193 B.

L8: Entry 5 of 10

File: USPT

Jan 8, 2002

US-PAT-NO: 6337195

DOCUMENT-IDENTIFIER: US 6337195 B1

** See image for Certificate of Correction **

TITLE: Colon specific genes and proteins

DATE-ISSUED: January 8, 2002

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Yu; Guo-Liang Darnestown MD
Rosen; Craiq Laytonsville MD

US-CL-CURRENT: 435/70.1; 530/350, 536/22.1

ABSTRACT:

Human colon specific gene polypeptides and DNA (RNA) encoding such polypeptides and a procedure for producing such polypeptides by recombinant techniques is disclosed. Also disclosed are methods for utilizing such polynucleotides or polypeptides as a diagnostic marker for colon cancer and as an agent to determine if colon cancer has metastasized. Also disclosed are antibodies specific to the colon specific gene polypeptides which may be used to target cancer cells and be used as part of a colon cancer vaccine. Methods of screening for agonists and antagonists for the polypeptide and therapeutic uses of the antagonists are disclosed.

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11 Claims, 17 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 17

6. Document ID: US 6312937 B1

L8: Entry 6 of 10

File: USPT

Nov 6, 2001

US-PAT-NO: 6312937

DOCUMENT-IDENTIFIER: US 6312937 B1

TITLE: Metalloproteinases

DATE-ISSUED: November 6, 2001

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP	CODE	COUNTRY
Ni; Jian	Rockville	MD			
Ruben; Steve	Olney	MD			
Brewer; Laurie	Poolesville	MD			
Gentz; Reiner	Silver Spring	MD a			
Rosen; Craiq	Laytonsville	MD			

US-CL-CURRENT: 435/219; 435/212, 435/226

ABSTRACT:

The present invention relates to novel metalloproteinase—like proteins. In particular, isolated nucleic acid molecules are provided encoding the human TACE—like and matrilysin—like proteins. TACE—like and matrilysin—like polypeptides are also provided as are vectors, host cells and recombinant methods for producing the same. The invention further relates to screening methods for identifying agonists and antagonists of TACE—like and matrilysin—like activity. Also provided are diagnostic methods for detecting cancer and therapeutic methods for cancer and other disorders characterized by an over or under production of these metalloproteinases.

30 Claims, 9 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 9

Claims KWC Dray	Reference				Front	Citation	Title	Full
Claims KWC Draw	Reference	Date	Classification	Review	Frent	Citation	2	Title

7. Document ID: US 6251648 B1

L8: Entry 7 of 10

File: USPT

Jun 26, 2001

US-PAT-NO: 6251648

DOCUMENT-IDENTIFIER: US 6251648 B1

TITLE: Gene encoding human Dnase

DATE-ISSUED: June 26, 2001

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Rosen; Craig Laytonsville MD

h eb bgeeef eheh ef b

Ruben; Steven M.

Olney

MD

Adams; Mark D.

North Potomac

MD

US-CL-CURRENT: 435/199; 530/300, 530/324

ABSTRACT:

A human DNase polypeptide and DNA (RNA) encoding such polypeptide and a procedure for producing such polypeptide by recombinant techniques is disclosed. Also disclosed are methods for utilizing such polypeptide for preventing and/or treating bronchopulmonary conditions. Diagnostic assays for identifying mutations in nucleic acid sequence encoding a polypeptide of the present invention and for detecting altered levels of the polypeptide of the present invention are also disclosed.

51 Claims, 7 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 7

Full Title Citation Front	Review Classification	Date Reference	Glaims KMC	Drawi Des
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8. Document ID:	US 6046031 A			
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US-PAT-NO: 6046031

DOCUMENT-IDENTIFIER: US 6046031 A

** See image for Certificate of Correction **

TITLE: Metalloproteinases

DATE-ISSUED: April 4, 2000

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY Ni; Jian Rockville MD Ruben; Steve Olney MD Brewer; Laurie Poolesville MD Gentz; Reiner Silver Spring MD Rosen; Craig Laytonsville MD

US-CL-CURRENT: $\underline{435/69.1}$; $\underline{435/219}$, $\underline{435/226}$, $\underline{435/252.33}$, $\underline{435/320.1}$, $\underline{435/325}$, $\underline{435/69.3}$, <u>536/23.1</u>, <u>536/23.2</u>, <u>536/23.5</u>

ABSTRACT:

The present invention relates to novel metalloproteinase-like proteins. In particular, isolated nucleic acid molecules are provided encoding the human TACE-like and matrilysin-like proteins. TACE-like and matrilysin-like polypeptides are also provided as are vectors, host cells and recombinant methods for producing the same. The invention further relates to screening methods for identifying agonists and antagonists of TACE-like and matrilysin-like activity. Also provided are diagnostic methods for detecting cancer and therapeutic methods for cancer and other disorders characterized by an over or under production of these metalloproteinases.

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60 Claims, 9 Drawing figures Exemplary Claim Number: 1

Number of Drawing Sheets: 9

Full Title Citation Front Review Classification Date Reference Claims KMC Draw Desc

9. Document ID: US 5733748 A

L8: Entry 9 of 10

File: USPT

Mar 31, 1998

US-PAT-NO: 5733748

DOCUMENT-IDENTIFIER: US 5733748 A

TITLE: Colon specific genes and proteins

DATE-ISSUED: March 31, 1998

INVENTOR-INFORMATION:

NAME CITY

STATE

ZIP CODE COUNTRY

Darnestown

Yu; Guo-Liang Rosen; Craiq

Laytonsville

MD MD

US-CL-CURRENT: 435/70.1; 435/252.3, 435/320.1, 435/325, 536/22.1, 536/23.5

ABSTRACT:

Human colon specific gene polypeptides and DNA (RNA) encoding such polypeptides and a procedure for producing such polypeptides by recombinant techniques is disclosed. Also disclosed are methods for utilizing such polynucleotides or polypeptides as a diagnostic marker for colon cancer and as an agent to determine if colon cancer has metastasized. Also disclosed are antibodies specific to the colon specific gene polypeptides which may be used to target cancer cells and be used as part of a colon cancer vaccine. Methods of screening for agonists and antagonists for the polypeptide and therapeutic uses of the antagonists are disclosed.

20 Claims, 17 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 17

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		Challen Freed Decision Change and a Decision Decision	Citation Front Review Classification Date Reference	Citation Front Review Classification Date Reference	Citation Front Review Classification Date Reference Claims KMC

10. Document ID: WO 9519985 A1

L8: Entry 10 of 10

File: EPAB

Jul 27, 1995

PUB-NO: WO009519985A1

DOCUMENT-IDENTIFIER: WO 9519985 A1 TITLE: HAEMOPOIETIC MATURATION FACTOR

PUBN-DATE: July 27, 1995

INVENTOR-INFORMATION:

NAME

COUNTRY

KIRKNESS, EWEN ADAMS, MARK D

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OLSEN, HENRIK ROSEN, CRAIG

 $\text{INT-CL (IPC): } \underline{\text{C07}} \ \underline{\text{H}} \ \underline{17/00}; \ \underline{\text{C12}} \ \underline{\text{N}} \ \underline{15/00}; \ \underline{\text{C12}} \ \underline{\text{N}} \ \underline{1/20}; \ \underline{\text{C12}} \ \underline{\text{P}} \ \underline{21/06}; \ \underline{\text{C07}} \ \underline{\text{K}} \ \underline{13/00}; \ \underline{\text{A61}} \ \underline{\text{K}}$

<u>37/00</u>

EUR-CL (EPC): C07K014/475

ABSTRACT:

Disclosed is a human maturation factor polypeptide and DNA(RNA) encoding such haemopoietic maturation factor polypeptides. Also provided is a procedure for producing such polypeptide by recombinant techniques and antibodies against such polypeptide. Such polypeptides may be combined with a suitable pharmaceutical carrier or diluent to provide diagnostic, therapeutic and/or prophylactic effects against various diseases related to the underexpression of such human haemopoietic maturation factor polypeptide.

Full Title Citation Front Review	Classification Date Reference	Claims KMC Draw. Des
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Terms	Documents	
Rosen-Craig.IN.		10

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Search Results - Record(s) 1 through 19 of 19 returned.

1: Document ID: AU 2003219999 A1, US 20030175340 A1, WO 2003075884 A1, WO 2004035004 A2

Using default format because multiple data bases are involved.

L10: Entry 1 of 19

File: DWPI

Sep 22, 2003

DERWENT-ACC-NO: 2004-020713

DERWENT-WEEK: 200431

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TITLE: Effervescent composition useful for treating osteoporosis comprises a bisphosphonate, an acid component and an alkaline effervescing component

INVENTOR: MCCALLISTER, D; ROSEN, C

PRIORITY-DATA: 2002US-0273081 (October 17, 2002), 2002US-0092083 (March 6, 2002)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
AU 2003219999 A1	September 22, 2003	•	000	A61K009/00
US 20030175340 A1	September 18, 2003		013	A61K031/675
WO 2003075884 A1	September 18, 2003	E	000	A61K009/00
WO 2004035004 A2	April 29, 2004	E	000	A61K000/00

INT-CL (IPC): $\underline{A61}$ \underline{K} $\underline{0/00}$; $\underline{A61}$ \underline{K} $\underline{9/00}$; $\underline{A61}$ \underline{K} $\underline{9/46}$; $\underline{A61}$ \underline{K} $\underline{31/4439}$; $\underline{A61}$ \underline{K} $\underline{31/66}$; $\underline{A61}$ \underline{K} $\underline{31/663}$; $\underline{A61}$ \underline{K} $\underline{31/675}$

Full	Title	Citation	Front Revie	ew Classification	Date Re	ference				Claims	KWIC	Draw.	Des
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	2.	Documer	nt ID: US	20020146778	A1								
L10:	Entr	y 2 of 1	.9		F	ile:	DWPI			Oct	10,	2002	

DERWENT-ACC-NO: 2003-255127

DERWENT-WEEK: 200325

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TITLE: Isolated polynucleotide encoding pineal gland specific gene-1 protein (PGSG-1), useful for regulation of the pituitary gland and for modulating biological rhythms

INVENTOR: HE, W W; ROSEN, C

PRIORITY-DATA: 1995US-0461248 (June 5, 1995), 2002US-0153739 (May 24, 2002)

PATENT-FAMILY:

PUB-NO PUB-DATE LANGUAGE PAGES MAIN-IPC

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021 C12P021/02

INT-CL (IPC): <u>C07 H 21/04</u>; <u>C07 K 14/575</u>; <u>C12 N 5/06</u>; <u>C12 P 21/02</u>

ABSTRACTED-PUB-NO: US20020146778A

BASIC-ABSTRACT:

NOVELTY - A new isolated polynucleotide (I) comprises:

- (a) a sequence encoding a polypeptide of 345 amino acids, fully defined in the specification;
- (b) a sequence encoding amino acid 22 to 283 of (a);
- (c) a sequence capable of hybridizing to and which is at least 70% identical to (a) or (b);
- (d) the DNA contained in ATCC Deposit No. 97162; or
- (e) a fragment of (a), (b), or (c).

DETAILED DESCRIPTION - INDEPENDENT CLAIMS are included for the following:

- A vector containing (I);
- (2) A host cell genetically engineered with the vector;
- (3) Producing a polypeptide comprising expressing from the host cell the polypeptide encoded by the DNA;
- (4) Producing cells capable of expressing a polypeptide comprising transforming or transfecting the cells with the vector;
- (5) A polypeptide (II) comprising a sequence of 345 amino acids, fully defined in the specification, amino acids 22 to 283 of (II), or a polypeptide encoded by the cDNA of ATCC Deposit No. 97162 and its fragments;
- (6) A compound effective as agonist, or antagonist for the polypeptide; and
- (7) Identifying compounds which bind to and activate or inhibit a receptor for the polypeptide comprising:
- (a) contacting a cell expressing on its surface a receptor for the polypeptide, the receptor being associated with a second component capable of providing a detectable signal in response to the binding of a compound to the receptor, with a compound to be screened under conditions to permit binding to the receptor; and
- (b) determining whether the compound binds to and activates or inhibits the receptor by detecting the presence or absence of a signal generated from the interaction of the compound with the receptor.

ACTIVITY - Endocrine; Cytostatic; Anticonvulsant; Ophthalmological.

No biological data given.

MECHANISM OF ACTION - Gene Therapy.

No biological data given.

USE - The methods are useful for diagnosing a disease or a susceptibility to a disease related to expression of the polypeptide comprising determining a mutation in the nucleic acid sequence encoding the polypeptide, treating a patient in need of pineal gland specific gene-1 (PGSG-1) comprising administering to the patient an

Page 3 of 9

amount of the polypeptide, and for diagnosis comprising analyzing for the presence of the polypeptide in a sample derived from a host (all claimed). The polypeptides are used for the regulation of the pituitary gland and to modulate biological rhythms. PGSG-1 is useful for treating conditions resulting from pineal gland tumors such as precocious puberty, hydrocephalus, papilledema, paralysis of upward gaze, ptosis and loss of pupillary light and accommodation reflexes.

3. Document ID: WO 200273366 A2, US 20030050070 A1, AU 2002254215 A1

L10: Entry 3 of 19

File: DWPI

Sep 19, 2002

DERWENT-ACC-NO: 2002-643728

DERWENT-WEEK: 200435

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TITLE: Method of dynamically allocating spectrum bandwidth by detecting criteria data sets for respective carriers and transmitting requests for a switch of carriers and transceivers over a control channel

INVENTOR: MASHINSKY, A; ROSEN, C; MASHINSKY,

PRIORITY-DATA: 2002US-357545P (February 15, 2002), 2001US-275818P (March 14, 2001), 2002US-0099552 (March 14, 2002)

PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
WO 200273366 A2	September 19, 2002	E	058	G06F000/00
US 20030050070 A1	March 13, 2003		000	H04Q007/20
AU 2002254215 A1	September 24, 2002		000	G06F000/00

INT-CL (IPC): $\underline{G06} \ \underline{F} \ \underline{0/00}; \ \underline{H04} \ \underline{B} \ \underline{1/38}; \ \underline{H04} \ \underline{M} \ \underline{1/00}; \ \underline{H04} \ \underline{Q} \ \underline{7/20}$

ABSTRACTED-PUB-NO: WO 200273366A

BASIC-ABSTRACT:

NOVELTY - Spectrum and network availability and congestion information from different service providers is pooled in a central database. Wholesale volumes of network capacity or accounts can be purchased and dynamically allocated to devices of different origin and ownership. A central system administrator re-bills and reconciles fractional usage to each device. Emergency calls can be given high priority to ensure their connection.

DETAILED DESCRIPTION - INDEPENDENT CLAIMS are included for

- (a) a device for dynamically switching communication modes in a wireless network
- (b) and a system for managing available spectrum in a wireless network with two or more available carriers

USE - Spectrum allocation in wireless telephone data systems.

ADVANTAGE - Efficient management of a network, especially at times of high congestion.

Full | Title | Citation | Front | Review | Classification | Date | Reference | Claims | Claims | KMC | Draw Desc

4. Document ID: US 20020086314 A1

L10: Entry 4 of 19

File: DWPI

Jul 4, 2002

DERWENT-ACC-NO: 2002-635672

DERWENT-WEEK: 200455

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TITLE: Novel human colon specific gene polypeptide, useful for treating colon cancer,

and as a diagnostic marker for colon cancer or matastasis of colon cancers

INVENTOR: ROSEN, C ; YU, G

PRIORITY-DATA: 1995US-0469667 (June 6, 1995), 1998US-0224110 (March 31, 1998),

2001US-0988292 (November 19, 2001)

PATENT-FAMILY:

PUB-NO

PUB-DATE

LANGUAGE

PAGES

MAIN-IPC

US 20020086314 A1

July 4, 2002

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C12Q001/68

INT-CL (IPC): $\underline{\text{C07}}$ $\underline{\text{H}}$ $\underline{\text{21/04}}$; $\underline{\text{C12}}$ $\underline{\text{N}}$ $\underline{\text{5/06}}$; $\underline{\text{C12}}$ $\underline{\text{N}}$ $\underline{\text{9/16}}$; $\underline{\text{C12}}$ $\underline{\text{P}}$ $\underline{\text{21/02}}$; $\underline{\text{C12}}$ $\underline{\text{Q}}$ $\underline{\text{1/68}}$; $\underline{\text{G01}}$ $\underline{\text{N}}$ $\underline{\text{33/574}}$

ABSTRACTED-PUB-NO: US20020086314A

BASIC-ABSTRACT:

NOVELTY - A human colon specific gene (CSG) polypeptide (I) comprising a sequence (S1) of 323 amino acids given in the specification, or its fragment, analog or derivative, or a sequence encoded by a human gene having a coding portion whose DNA has at least 90% identity to a sequence (S2) of 638, 1209, 548, 878, 560, 709, 559, 409 or 600 base pairs given in the specification, is new.

DETAILED DESCRIPTION - A human colon specific polypeptide (I) comprising a sequence (S1) of 323 amino acids fully defined in the specification, or its fragment, analog or derivative, or a sequence encoded by a human gene having a coding portion whose DNA has at least 90% identity to a sequence (S2) of 638, 1209, 548, 878, 560, 709, 559, 409 or 600 base pairs fully defined in the specification, and a polypeptide encoded by the human gene whose coding region includes a DNA having at least 90% identity to the DNA contained in ATCC 97102 and fragments, analogs or derivatives of the polypeptide.

INDEPENDENT CLAIMS are also included for:

- (1) an isolated polynucleotide (II) comprising:
- (a) a polynucleotide encoding S1
- (b) a polynucleotide capable of hybridizing to and which is at least 70% identical to (a): and
- (c) a polynucleotide encoding the same mature polypeptide as a human gene having a coding portion which includes DNA having at least 90% identity to S2 or to the DNA included in ATCC 97102;
- (2) a vector (III) containing (II);
- (3) a host cell (IV) transformed or transfected with (III);
- (4) production of (I);

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(5) producing cells capable of expressing (I) by genetically engineering cells with (III);

- (6) an antibody (V) against (I);
- (7) a compound (VI) which inhibits activation of (I);
- (8) diagnosing (M) a disorder of colon in a patient by determining transcription of a human gene in a sample derived from a non-colon tissue of a host (the gene has a coding portion which includes a DNA having at least 90% identity to a DNA selected from S2 or a sequence of 874, 570, 1121 or 605 base pairs fully defined in the specification);
- (9) an isolated antibody (VII) or its portion that specifically binds to a protein
- (P) or is produced by immunizing an animal with (P) ((P) is:
- (a) a protein whose sequence consists of amino acid residues 1-323 of S1, or a protein consisting of a fragment comprising at least 30 or 50 contiguous residues of S1 ((VII) or its portion specifically binds to S1); and
- (b) a protein whose sequence consists of the amino acid sequence of the full-length or mature CSG10 polypeptide encoded by the cDNA contained in ATCC 97102, or a protein consisting of a fragment of the CSG10 polypeptide encoded by the cDNA contained in ATCC 97102 (the fragment comprises at least 30 or 50 amino acid residues of the CSG10 polypeptide encoded by the cDNA contained in ATCC 97102 and (VII) or its portion binds specifically to the CSG10 polypeptide encoded by the cDNA contained in ATCC 97102));

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- (10) an isolated cell (VIII) that produces (VII); and
- (11) a hybridoma (IX) that produces (VII).

ACTIVITY - Cytostatic.

MECHANISM OF ACTION - Inhibitor of activation of (I) (claimed); gene therapy; vaccine.

No supporting data is given.

USE - (I) Is useful for treating a patient having need of (I).

- (VI) Is useful for the treatment of a patient and inhibit expression and activity (I). (VI) Is a polypeptide and a therapeutically effective amount of the compound is administered by providing to the patient DNA encoding the polypeptide and expressing the polypeptide in vivo.
- (VII) Is useful for detecting CSG10 protein in a biological sample by contacting the biological sample with (VII) or its portion, and detecting the CSG10 protein in the biological sample (claimed).
- (I) Or (II) is useful as a diagnostic marker for colon cancer, or as an agent to determine if colon cancer has metastasized, and for in vitro purposes related to scientific research, synthesis of DNA and manufacture of DNA vectors.
- (V) Is useful to target cancer cells and as a part of a colon cancer vaccine.
- (I) Is useful for treating colon cancer, to screen for compounds which interact with (I), for example, compounds which inhibit or activate (I), and as an immunogen to produce antibodies to (I). (II) Is useful in gene therapy and for chromosome identification.

5. Document ID: US 6337195 B1

L10: Entry 5 of 19 File: DWPI Jan 8, 2002

DERWENT-ACC-NO: 2002-163239

DERWENT-WEEK: 200455

COPYRIGHT 2004 DERWENT INFORMATION LTD

TITLE: Human colon specific gene polypeptide, useful as diagnostic marker, vaccine

and for screening agonists and antagonists for treating colon cancer

INVENTOR: ROSEN, C; YU, G

PRIORITY-DATA: 1995US-0469667 (June 6, 1995), 1998US-0224110 (March 31, 1998)

PATENT-FAMILY:

 PUB-NO
 PUB-DATE
 LANGUAGE
 PAGES
 MAIN-IPC

 US 6337195 B1
 January 8, 2002
 049
 C07H021/04

INT-CL (IPC): <u>C07</u> <u>H</u> <u>21/04</u>; <u>C07</u> <u>K</u> <u>13/00</u>; <u>C12</u> <u>P</u> <u>21/04</u>

ABSTRACTED-PUB-NO: US 6337195B

BASIC-ABSTRACT:

NOVELTY - An isolated human colon specific gene polypeptide (I) comprising a sequence (S1) 95% identical to a sequence (S2) consisting of amino acids 2-323 of a fully defined sequence (S3) of 323 amino acids as given in the specification, where (I) binds to an antibody that specifically binds to a protein consisting of (S3), is new.

DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for the following:

- (1) preparation of (I); and
- (2) a polypeptide (II) comprising at least 30 contiguous amino acids of (S3).

ACTIVITY - Cytostatic.

MECHANISM OF ACTION - Vaccine. No supporting data is given.

USE - (I) is useful as diagnostic marker for colon cancer and as a colon cancer vaccine. It is also useful for targeting cancer cells and for screening agonist and antagonist for (I) which are useful for treating colon cancer.

	Full	Title	Citation	Fient	Review	Classification	Date	Referenc	e l		Claims	KMC	Draw Desc
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		6.	Docume	ent ID	: US 6	175831 B1	•						
I	10:	Entr	y 6 of	19				File:	DWPI		Jan	16,	2001

DERWENT-ACC-NO: 2001-463379

DERWENT-WEEK: 200150

COPYRIGHT 2004 DERWENT INFORMATION LTD

TITLE: Network database system has database server which is responsive to parser processing to manipulate record in database and selected records are linked by confirmed defined relationship

INVENTOR: BERLYN, N D; BODDU, C; CHIBNIK, R; CLIFFORD, S; GREEN, J; HABER, D; MITCHELL, L; ROSEN, C; SALAMON, M R; SAMUELS, D; SEIFER, A; WEINREICH, A P; ZILBERBERG, S

PRIORITY-DATA: 1997US-0785559 (January 17, 1997)

PATENT-FAMILY:

PUB-NO PUB-DATE LANGUAGE PAGES MAIN-IPC
US 6175831 B1 January 16, 2001 048 G06F017/30

INT-CL (IPC):  $\underline{606} + \underline{17/30}$ 

ABSTRACTED-PUB-NO: US 6175831B

BASIC-ABSTRACT:

NOVELTY - Database connectivity engine pre-processing output of web server, is coupled to database server to which queue watcher is coupled. Mail server coupled to communication port to receive incoming e-mails, is coupled to watcher to transmit outgoing e-mails. Parser coupled to mail server processes incoming e-mails, is coupled to database server that manipulates a record. Selected records are linked by confirmed defined relationship.

DETAILED DESCRIPTION - The web server is connected to communication port. A database server is connected to the database which has a number records. An INDEPENDENT CLAIM is also included for a method of creating a networking database system.

USE - Networking database system.

ADVANTAGE - The networking database has applications for searching in terms of finding other individuals in the database, finding a connection to other users in the database. The system finds other individuals in the database having particular professional or personal characteristics or features that are of interest to other members. Thus the system performs search using the database and the defined relationships in order to determined specific information about registered user.

DESCRIPTION OF DRAWING(S) - The figure shows the flowchart of illustrating the process, add new relationship to a personal profile of networking database system.

Full	Title	Citation	Frent	Review	Classification	Date	Reference	Claims	Kwic	Drawt Desc

# 7. Document ID: WO 9854963 A2, AU 9878120 A, EP 1039801 A1, JP 2002516573 W, US 20030092893 A1, EP 1428833 A2

L10: Entry 7 of 19

File: DWPI

Dec 10, 1998

DERWENT-ACC-NO: 1999-059865

DERWENT-WEEK: 200462

COPYRIGHT 2004 DERWENT INFORMATION LTD

TITLE: New isolated human genes and the secreted polypeptides they encode - useful for diagnosis and treatment of e.g. cancers, neurological disorders, immune diseases, inflammation or blood disorders

INVENTOR: BREWER, L A; CARTER, K; DILLON, P; EBNER, R; ENDRESS, G; FAN, P; FENG, P; FERRIE, A M; FISCHER, C; FLORENCE, C; FLORENCE, K; GREENE, J; HU, J; KYAW, H; LAFLEUR, D; LI, Y; MOORE, P A; NI, J; OLSEN, H; ROSEN, C; RUBEN, S; SHI, Y E; SOPPET, D; WEI, Y; YOUNG, P; YU, G; ZENG, Z; CARTER, K C; DILLON, P J; ENDRESS, G A; FISCHER, C L; GREENE, J M; LAFLEUR, D W; MORE, P A; OLSEN, H S; ROSEN, C A; RUBEN, S M; SHI, Y; SOPPET, D R

Record List Display Page 8 of 9

PRIORITY-DATA: 1997US-070923P (December 18, 1997), 1997US-048875P (June 6, 1997), 1997US-048876P (June 6, 1997), 1997US-048877P (June 6, 1997), 1997US-048878P (June 6, 1997), 1997US-048880P (June 6, 1997), 1997US-048881P (June 6, 1997), 1997US-048882P (June 6, 1997), 1997US-048883P (June 6, 1997), 1997US-048884P (June 6, 1997), 1997US-048885P (June 6, 1997), 1997US-048892P (June 6, 1997), 1997US-048893P (June 6, 1997), 1997US-048894P (June 6, 1997), 1997US-048895P (June 6, 1997), 1997US-048896P (June 6, 1997), 1997US-048897P (June 6, 1997), 1997US-048898P (June 6, 1997), 1997US-048899P (June 6, 1997), 1997US-048900P (June 6, 1997), 1997US-048901P (June 6, 1997), 1997US-048915P (June 6, 1997), 1997US-048916P (June 6, 1997), 1997US-048917P (June 6, 1997), 1997US-048949P (June 6, 1997), 1997US-048962P (June 6, 1997), 1997US-048963P (June 6, 1997), 1997US-048964P (June 6, 1997), 1997US-048970P (June 6, 1997), 1997US-048971P (June 6, 1997), 1997US-048972P (June 6, 1997), 1997US-048974P (June 6, 1997), 1997US-049019P (June 6, 1997), 1997US-049020P (June 6, 1997), 1997US-049373P (June 6, 1997), 1997US-049374P (June 6, 1997), 1997US-049375P (June 6, 1997), 1997US-057584P (September 5, 1997), 1997US-057627P (September 5, 1997), 1997US-057628P (September 5, 1997), 1997US-057629P (September 5, 1997), 1997US-057634P (September 5, 1997), 1997US-057635P (September 5, 1997), 1997US-057642P (September 5, 1997), 1997US-057643P (September 5, 1997), 1997US-057644P (September 5, 1997), 1997US-057645P (September 5, 1997), 1997US-057646P (September 5, 1997), 1997US-057647P (September 5, 1997), 1997US-057648P (September 5, 1997), 1997US-057649P (September 5, 1997), 1997US-057650P (September 5, 1997), 1997US-057651P (September 5, 1997), 1997US-057654P (September 5, 1997), 1997US-057661P (September 5, 1997), 1997US-057662P (September 5, 1997), 1997US-057666P (September 5, 1997), 1997US-057667P (September 5, 1997), 1997US-057668P (September 5, 1997), 1997US-057760P (September 5, 1997), 1997US-057761P (September 5, 1997), 1997US-057762P (September 5, 1997), 1997US-057763P (September 5, 1997), 1997US-057764P (September 5, 1997), 1997US-057765P (September 5, 1997), 1997US-057769P (September 5, 1997), 1997US-057770P (September 5, 1997), 1997US-057771P (September 5, 1997), 1997US-057774P (September 5, 1997), 1997US-057775P (September 5, 1997), 1997US-057776P (September 5, 1997), 1997US-057777P (September 5, 1997), 1997US-057778P (September 5, 1997), 1997US-049896P (June 6, 1997), 1998US-092921P (July 15, 1998), 1998US-094657P (July 30, 1998), 1998US-0205258 (December 4, 1998), 2001US-0023282 (December 20, 2001)

### PATENT-FAMILY:

PUB-NO	PUB-DATE	LANGUAGE	PAGES	MAIN-IPC
WO 9854963 A2	December 10, 1998	E	770	A01N037/18
AU 9878120 A	December 21, 1998		000	
EP 1039801 A1	October 4, 2000	E	000	A01N037/18
JP 2002516573 W	June 4, 2002		914	C12N015/09
US 20030092893 A1	May 15, 2003		000	C07K016/00
EP 1428833 A2	June 16, 2004	E	000	C07K014/435

INT-CL (IPC): A01 N 37/18; A01 N 43/04; A61 K 31/711; A61 K 38/00; A61 K 38/17; A61 K 39/395; A61 K 48/00; A61 P 7/00; A61 P 25/00; A61 P 29/00; A61 P 35/00; A61 P 37/00; A61 P 43/00; C07 K 14/435; C07 K 14/47; C07 K 16/00; C07 K 16/18; C12 N 1/15; C12 N 1/19; C12 N 1/20; C12 N 1/21; C12 N 5/00; C12 N 5/06; C12 N 5/10; C12 N 15/00; C12 N 1/02; C12 Q 1/02; C12 Q 1/68; G01 N 33/53

ABSTRACTED-PUB-NO: WO 9854963A

BASIC-ABSTRACT:

An isolated nucleic acid molecule (NAM) (I) comprising a polynucleotide (PN) having a nucleotide sequence (NS) at least 95% identical to: (a) a PN fragment of one of a total of 207 defined human cDNA sequences given in the specification or a PN fragment of the cDNA sequence included in ATCC Deposit No. Z which is hybridisable to one of the 207 defined cDNA sequence; (b) a PN which is an (allelic) variant of one of the 207 defined cDNA sequences; (c) a PN encoding a biologically active polypeptide or a polypeptide fragment, domain or epitope of one of the 207 defined amino acid sequences given in the specification or a polypeptide fragment encoded by a cDNA sequence included in ATCC Deposit No. Z which is hybridisable to one of the defined

cDNA sequences; (d) a PN which encodes a species homologue of one of the 207 defined polypeptides; or (e) a PN capable of hybridising under stringent conditions to any one of the PNs specified in (a)-(d), where the PN

ef

# **Hit List**

Clear Generate Collection Print Fwd Refs Bkwd Refs Generate OACS

# Search Results - Record(s) 1 through 78 of 78 returned.

# 1. Document ID: US 20040132087 A1

# Using default format because multiple data bases are involved.

L13: Entry 1 of 78

File: PGPB

Jul 8, 2004

PGPUB-DOCUMENT-NUMBER: 20040132087

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040132087 A1

TITLE: Novel human enzyme family members and uses thereof

PUBLICATION-DATE: July 8, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Meyers, Rachel E. Newton MA US
Glucksmann, Maria Alexandria Lexington MA US
Rudolph-Owen, Laura A. Medford MA US

US-CL-CURRENT:  $\underline{435/6}$ ;  $\underline{435/226}$ ,  $\underline{435/320.1}$ ,  $\underline{435/325}$ ,  $\underline{435/69.1}$ ,  $\underline{530/350}$ ,  $\underline{536/23.2}$ 

Full Title Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	~	Draw, Desi
			-						

# 2. Document ID: US 20040121956 A1

L13: Entry 2 of 78

File: PGPB

Jun 24, 2004

PGPUB-DOCUMENT-NUMBER: 20040121956

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040121956 A1

TITLE: Drosophila G protein coupled receptors, nucleic acids, and methods related to

the same

PUBLICATION-DATE: June 24, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47 Lowery, David E. Portage MI US Smith, Valdin G. Kalamazoo MI US Kubiak, Teresa M. Richland ΜT US Larsen, Martha J. Kalamazoo MI US

US-CL-CURRENT: 514/12; 435/320.1, 435/348, 435/69.1, 530/350, 536/23.5

## ABSTRACT:

h e b b g e e e f e h eh ef b

The present invention provides a Drosophila melanogaster GPCR (DmGPCR) polypeptides and polynucleotides which identify and encode such a DmGPCR. In addition, the invention provides expression vectors, host cells and methods for its production. The invention also provides methods for the identification of homologs in other animals, and of DmGPCR agonists/antagonists, useful for the treatment of diseases in animals and for the control of insects that are injurious or harmful to plants or animals.

# 3. Document ID: US 20040110185 A1

L13: Entry 3 of 78

File: PGPB

Jun 10, 2004

PGPUB-DOCUMENT-NUMBER: 20040110185

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040110185 A1

TITLE: Human hypothalmic ("HR") receptor polypeptide compositions, methods and uses

thereof

PUBLICATION-DATE: June 10, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Duhl, David Oakland CA US

US-CL-CURRENT:  $\underline{435/6}$ ;  $\underline{435/320.1}$ ,  $\underline{435/325}$ ,  $\underline{435/69.1}$ ,  $\underline{530/350}$ ,  $\underline{536/23.5}$ 

### ABSTRACT:

A new human hypothalmic receptor has been identified, and the amino acid and nucleotide sequence of the receptor are provided. The nucleotide sequence is useful to construct expression cassettes and vectors to produce host cells which are capable of expressing the receptor, its mutants, fragments, or fusions. Such polypeptides are useful for identifying new agonists and antagonists.

"Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	******	KWC	Draw Des
							7-0-0					
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# 4. Document ID: US 20040110170 A1

L13: Entry 4 of 78

File: PGPB

Jun 10, 2004

PGPUB-DOCUMENT-NUMBER: 20040110170

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040110170 A1

TITLE: Cloning and characterization of calcitonin gene related peptide receptors

PUBLICATION-DATE: June 10, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Pisegna, Joseph R. Santa Monica CA US Wank, Stephen A. Potomac MD US

h eb bgeeef eheh ef be

US-CL-CURRENT:  $\underline{435/6}$ ;  $\underline{435/320.1}$ ,  $\underline{435/325}$ ,  $\underline{435/69.1}$ ,  $\underline{530/350}$ ,  $\underline{536/23.5}$ 

### ABSTRACT:

This invention provides CGRP receptors (including both amino acid and nucleic acid sequences). Compositions which include these polypeptides, proteins, nucleic acids, recombinant cells, transgenic animals, and antibodies to the receptors are also provided.

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	4	KOMIC	Draw. Desc
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5. Document ID: US 20040009553 A1

L13: Entry 5 of 78

File: PGPB

Jan 15, 2004

PGPUB-DOCUMENT-NUMBER: 20040009553

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040009553 A1

TITLE: Novel 27411, 23413, 22438, 23553, 25278, 26212, NARC SC1, NARC 10A, NARC 1, NARC 12, NARC 13, NARC17, NARC 25, NARC 3, NARC 4, NARC 7, NARC 8, NARC 11, NARC 14A, NARC 15, NARC 16, NARC 19, NARC 20, NARC 26, NARC 27, NARC 28, NARC 30, NARC 5, NARC 6, NARC 9, NARC 10C, NARC 8B, NARC 9, NARC2A, NARC 16B, NARC 1C, NARC1A, NARC 25, 86604 and 32222 molecules and uses therefor

PUBLICATION-DATE: January 15, 2004

## INVENTOR-INFORMATION:

CITY	STATE	COUNTRY	RULE-47
Lexington	MA	US	Α.
Saugus	MA	US	
Newton	MA	US	
Medford	MA	US	
Chestnut Hill	MA	US	
Newton	MA	US	
Edison	NJ	US	
Somerville	MA	US	•
	Lexington Saugus Newton Medford Chestnut Hill Newton Edison	Lexington MA Saugus MA Newton MA Medford MA Chestnut Hill MA Newton MA Edison NJ	Lexington MA US Saugus MA US Newton MA US Medford MA US Chestnut Hill MA US Newton MA US Edison NJ US

US-CL-CURRENT: 435/69.1; 435/320.1, 435/325, 530/350, 536/23.5

## ABSTRACT:

The invention provides isolated nucleic acids molecules and proteins, designated 27411, 23413, 22438, 23553, 25278, 26212, NARC SC1, NARC 10A, NARC 1, NARC 12, NARC 13, NARC 17, NARC 25, NARC 3, NARC 4, NARC 7, NARC 8, NARC 11, NARC 14A, NARC 15, NARC 16, NARC 19, NARC 20, NARC 26, NARC 27, NARC 28, NARC 30, NARC 5, NARC 6, NARC 9, NARC 10C, NARC 8B, NARC 9, NARC2A, NARC 16B, NARC 1C, NARC 1A, NARC 25, 86604 and 32222 nucleic acid molecules and proteins. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing said nucleic acid molecules, host cells into which the expression vectors have been introduced, nonhuman transgenic animals in which a said genes have been introduced or disrupted, fusion proteins, antigenic peptides and antibodies to said proteins. Diagnostic and therapeutic methods utilizing compositions of the invention are also provided.

ef

Full Title Citation Front Review Classification Date Reference Sequences Attachments 6000 Draw Desc

# 6. Document ID: US 20030215860 A1

L13: Entry 6 of 78

File: PGPB

Nov 20, 2003

PGPUB-DOCUMENT-NUMBER: 20030215860

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030215860 A1

TITLE: Novel 18636, 2466, 43238, 1983, 52881, 2398, 45449, 50289, 52872 and 26908

molecules and uses therefor

PUBLICATION-DATE: November 20, 2003

### INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Glucksmann, Maria A.	Lexington	MA	US	
Silos-Santiago, Inmaculada	Del Mar	CA	us	
Carroll, Joseph M.	Cambridge	MA	us	
Galvin, Katherine M.	Jamaica Plain	MA	US	

US-CL-CURRENT: 435/6; 435/320.1, 435/325, 435/69.1, 530/350, 530/388.1, 536/23.1

### ABSTRACT:

The invention provides isolated nucleic acids molecules, designated 18636, 2466, 43238, 1983, 52881, 2398, 45449, 50289, 52872 and 26908 nucleic acid molecules. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 18636, 2466, 43238, 1983, 52881, 2398, 45449, 50289, 52872 and 26908 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 18636, 2466, 43238, 1983, 52881, 2398, 45449, 50289, 52872 or 26908 gene has been introduced or disrupted. The invention still further provides isolated 18636, 2466, 43238, 1983, 52881, 2398, 45449, 50289, 52872 or 26908 proteins, fusion proteins, antigenic peptides and anti-18636, 2466, 43238, 1983, 52881, 2398, 45449, 50289, 52872 or 26908 antibodies. Diagnostic and therapeutic methods utilizing compositions of the invention are also provided.

Full	Title Citation Front	Review Classification Date	Reference	Sequences	Attachments	····	MIC	Draw Desc
	7. Document ID:	US 20030187222 A1		·	······································	•••••••••	******	
L13:	Entry 7 of 78		File:	PGPB		Oct	2,	2003

PGPUB-DOCUMENT-NUMBER: 20030187222

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030187222 A1

TITLE: Novel galanin receptor

PUBLICATION-DATE: October 2, 2003

INVENTOR-INFORMATION:

h e b b g e e e f e h eh ef b e

NAME CITY STATE COUNTRY RULE-47

Shi-Hsiang, Shen Beaconsfield CA
Sultan, Ahmad Dorval CA
Wahlestedt, Claes Montreal CA
Walker, Philippe Montreal CA

US-CL-CURRENT:  $\underline{530}/\underline{350}$ ;  $\underline{435}/\underline{320.1}$ ,  $\underline{435}/\underline{353}$ ,  $\underline{435}/\underline{455}$ ,  $\underline{435}/\underline{69.1}$ ,  $\underline{435}/\underline{7.1}$ ,  $\underline{530}/\underline{388.22}$ ,

536/23.5

### ABSTRACT:

The present invention is directed to a novel receptor for galanin which has been designated as galanin receptor 2. The invention encompasses both the receptor protein as well as nucleic acids encoding the protein. In addition, the present invention is directed to methods and compositions which rely upon either GAL-R2 proteins or nucleic acids.

Full Title Citation Front	Review Classification	Date Reference	Sequences /	Attachments   •	·	KOMC	Draw, Desi

# 8. Document ID: US 20030175883 A1

L13: Entry 8 of 78

File: PGPB

Sep 18, 2003

PGPUB-DOCUMENT-NUMBER: 20030175883

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030175883 A1

TITLE: DNA encoding a mammalian LPA receptor and uses thereof

PUBLICATION-DATE: September 18, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Bard, Jonathan A Doylestown PA U

US-CL-CURRENT: 435/69.1; 435/320.1, 435/325, 530/350, 536/23.5

# ABSTRACT:

This invention provides an isolated nucleic acid encoding a mammalian LPA receptor, a purified mammalian LPA receptor, vectors comprising isolated nucleic acid encoding an mammalian LPA receptor, cells comprising such vectors, antibodies directed to a mammalian LPA receptor, nucleic acid probes useful for detecting nucleic acid encoding a mammalian LPA receptor, antisense oligonucleotides complementary to unique sequences of nucleic acid encoding mammalian LPA receptor, transgenic, nonhuman animals which express DNA encoding a normal or a mutant mammalian LPA receptor, methods of isolating an mammalian LPA receptor, methods of treating an abnormality that is linked to the activity of the mammalian LPA receptor, as well as methods of determining binding of compounds to mammalian LPA receptors.

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	,,	KMC	Draw, Desc

9. Document ID: US 20030162944 A1

h e b b g e e e f e h eh ef b e

L13: Entry 9 of 78 File: PGPB Aug 28, 2003

PGPUB-DOCUMENT-NUMBER: 20030162944

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030162944 A1

TITLE: Nucleic acid encoding neuropeptide Y/peptide YY (Y2) receptors and uses

thereof

PUBLICATION-DATE: August 28, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Gerald, Christophe	Ridgewood	NJ	US	
Walker, Mary W.	Elmwood Park	NJ	US	
Branchek, Theresa	Teaneck	NJ	US	
Weinshank, Richard L.	Teaneck	NJ	US	

US-CL-CURRENT: <u>530</u>/<u>350</u>; <u>435</u>/<u>320.1</u>, <u>435</u>/<u>325</u>, <u>435</u>/<u>69.1</u>, <u>536</u>/<u>23.5</u>

### ABSTRACT:

This invention provides isolated nucleic acid molecules encoding Y2 receptors, an isolated, purified Y2 receptor protein, vectors comprising isolated nucleic acid molecules encoding Y2 receptors, mammalian, insect, bacterial and yeast cells comprising such vectors, antibodies directed to the Y2 receptors, nucleic acid probes useful for detecting nucleic acid encoding Y2 receptors, antisense oligonucleotides complementary to unique sequences of a nucleic acid molecule which encodes a Y2 receptor, pharmaceutical compounds related to the Y2 receptors, and nonhuman transgenic animals which express nucleic acid encoding a normal or mutant Y2 receptor. This invention further provides methods for determining ligand binding, detecting expression, drug screening, and methods of treatment involving Y2 receptors.

Full	Title Citation Front Review	Classification Date Reference	Sequences Attachments	- KNOC Drawn Desi
	10. Document ID: US 20	0020140440 A 1	***************************************	***************************************
T.13:	Entry 10 of 78	7030146449 A1 File:	DCDB	Aug 7 2003 /

PGPUB-DOCUMENT-NUMBER: 20030148449

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030148449 A1

TITLE: G protein coupled receptor agonists and antagonists and methods of activating and inhibiting G protein coupled receptors using the same

PUBLICATION-DATE: August 7, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47 Kuliopulos, Athan Winchester MA US Covic, Lidija Somerville MA US

US-CL-CURRENT: 435/69.1; 435/320.1, 435/325, 514/12, 514/558, 530/350

h e b b g e e e f e h e h e f b e

### ABSTRACT:

The invention relates generally to G protein coupled receptors and in particular to agonists and antagonists of G protein receptors and methods of using the same.

Full Title Citation Front Review Classification Date Reference Sequences Attachments --- KMC Draw Desc

## 11. Document ID: US 20030138890 A1

L13: Entry 11 of 78

File: PGPB

Jul 24, 2003

PGPUB-DOCUMENT-NUMBER: 20030138890

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030138890 A1

TITLE: Novel G protein-coupled receptor family members, human thioredoxin family members, human leucine-rich repeat family members, and human ringfinger family member

PUBLICATION-DATE: July 24, 2003

#### INVENTOR-INFORMATION:

CITY	STATE	COUNTRY	RULE-47
Lexington	MA	US	
Jamaica Plain	MA	US	
Jamaica Plain	MA	US	
Brookline	MA	US	
Framingham	MA	US	
Watertown	MA	US	
Chestnut Hill	MA	US	•
	Lexington Jamaica Plain Jamaica Plain Brookline Framingham Watertown	Lexington MA  Jamaica Plain MA  Jamaica Plain MA  Brookline MA  Framingham MA  Watertown MA	Lexington MA US  Jamaica Plain MA US  Jamaica Plain MA US  Brookline MA US  Framingham MA US  Watertown MA US

US-CL-CURRENT:  $\underline{435}/\underline{69.1}$ ;  $\underline{435}/\underline{320.1}$ ,  $\underline{435}/\underline{325}$ ,  $\underline{530}/\underline{350}$ ,  $\underline{536}/\underline{23.5}$ 

### ABSTRACT:

The invention provides isolated nucleic acids molecules, designated 20716, 65494, 44576, 1983, 52881, 2398, 45449, 50289, 52872, 22105, 22109, 22108, 47916, 33395, 31939, and 84241 nucleic acid molecules, which encode novel G protein-coupled receptor family members, human thioredoxin family members, human leucine-rich repeat family members, and human ringfinger family member. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 20716, 65494, 44576, 1983, 52881, 2398, 45449, 50289, 52872, 22105, 22109, 22108, 47916, 33395, 31939, or 84241 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 20716, 65494, 44576, 1983, 52881, 2398, 45449, 50289, 52872, 22105, 22109, 22108, 47916, 33395, 31939, or 84241 gene has been introduced or disrupted. The invention still further provides isolated 20716, 65494, 44576, 1983, 52881, 2398, 45449, 50289, 52872, 22105, 22109, 22108, 47916, 33395, 31939, or 84241 proteins, fusion proteins, antigenic peptides and anti-20716, 65494, 44576, 1983, 52881, 2398, 45449, 50289, 52872, 22105, 22109, 22108, 47916, 33395, 31939, or 84241 antibodies. Diagnostic methods utilizing compositions of the invention are also provided.

Full	Title Citati	n Front	Review	Classification	Date	Reference	Sequences	Attachmenta	,,	KWIC	Draw Desc

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12. Document ID: US 20030119096 A1

L13: Entry 12 of 78

File: PGPB

Jun 26, 2003

PGPUB-DOCUMENT-NUMBER: 20030119096

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030119096 A1

TITLE: Method of treating an abnormality using a GALR3 receptor antagonist

PUBLICATION-DATE: June 26, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47 Bard, Jonathan A. Doylestown PA US Borowsky, Beth Montclair NJ US Smith, Kelli E. Fair Lawn NJ US

US-CL-CURRENT: 435/69.1; 435/320.1, 435/325, 530/350, 536/23.5

### ABSTRACT:

This invention provides an isolated nucleic acid encoding a mammalian galanin receptor, an isolated galanin receptor protein, vectors comprising isolated nucleic acid encoding a mammalian galanin receptor, cells comprising such vectors, antibodies directed to a mammalian galanin receptor, nucleic acid probes useful for detecting nucleic acid encoding a mammalian galanin receptor, antisense oligonucleotides complementary to unique sequences of nucleic acid encoding a mammalian galanin receptor, nonhuman transgenic animals which express DNA encoding a normal or a mutant mammalian galanin receptor, as well as methods of determining binding of compounds to mammalian galanin receptors.

Full Tit	le Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	,	 KOMC	Drawt Des
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13. Document ID: US 20030099970 A1

L13: Entry 13 of 78

File: PGPB

May 29, 2003

PGPUB-DOCUMENT-NUMBER: 20030099970

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030099970 A1

TITLE: Human-derived bradeion proteins, DNA coding for the proteins, and uses thereof

PUBLICATION-DATE: May 29, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Tanaka, Manami Ibaraki JP

Tanaka, Tomoo Kanagawa JP

US-CL-CURRENT: 435/6; 435/320.1, 435/325, 435/69.1, 435/7.23, 530/350, 530/388.22, 536/23.5

ABSTRACT:

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A human-derived bradeion protein, which has the following properties: (i) it is a transmembranous protein; (ii) it has a structure characteristic of growth hormone and cytokine receptors even in a structure of its transmembranous portion when its structure is determined by a hydrophobicity analysis according to Kyte-Doolittle method; (iii) it is expressed at a high level in a human adult brain, and in less amount in the heart, while it is not expressed in other adult organs or fetus; (iv) it induces programmed cell death (apoptosis) when over-expressed in a cultured human nerve cell lines; (v) it induces termination of cell division and aging when over-expressed in a cultured human normal cell; (vi) it is located in cytoplasm, and forms an intracellular aggregate when overexpressed; and (vii) besides human adult neurons, it is specifically expressed in a human colorectal cancer cell line or in a skin cancer cell line, or an analogue thereof.

Full Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	J	Drawt Desc
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14. Document ID: US 20030082738 A1

L13: Entry 14 of 78

File: PGPB

May 1, 2003

PGPUB-DOCUMENT-NUMBER: 20030082738

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030082738 A1

TITLE: 1983, 52881, 2398, 45449, 50289, and 52872, novel G protein-coupled receptors

and uses therefor

PUBLICATION-DATE: May 1, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47 Glucksmann, Maria Alexandra Lexington US Galvin, Katherine M. Jamaica Plain MA US Silos-Santiago, Inmaculada Cambridge MA US

US-CL-CURRENT: 435/69.1; 435/320.1, 435/325, 530/350, 536/23.5

## ABSTRACT:

The invention provides isolated nucleic acids molecules, designated 1983, 52881, 2398, 45449, 50289, and 52872 nucleic acid molecules, which encode novel G protein-coupled receptor members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 1983, 52881, 2398, 45449, 50289, or 52872 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 1983, 52881, 2398, 45449, 50289, or 52872 gene has been introduced or disrupted. The invention still further provides isolated 1983, 52881, 2398, 45449, 50289, or 52872 proteins, fusion proteins, antigenic peptides and anti-1983, 52881, 2398, 45449, 50289, or 52872 antibodies. Diagnostic methods utilizing compositions of the invention are also provided.

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Atlachments	,,	Draw Desc

15. Document ID: US 20030082641 A1

L13: Entry 15 of 78

File: PGPB

May 1, 2003

h e b b g e e e f e h eh ef b e

PGPUB-DOCUMENT-NUMBER: 20030082641

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030082641 A1

TITLE: A METHOD OF TREATING DEPRESSION USING A GALR3 RECEPTOR ANTAGONIST

PUBLICATION-DATE: May 1, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Bard, Jonathan A.	Doylestown	PA	US	
Borowsky, Beth	Montclair	NJ	US	
Smith, Kelli E.	Wayne	NJ	US	
Branchek, Theresa A.	Teaneck	NJ	US	
Gerald, Christophe P.G.	Ridgewood	NJ	US	
Jones, Kenneth A.	Bergenfield	NJ	US	

US-CL-CURRENT:  $\underline{435}/\underline{7.21}$ ;  $\underline{435}/\underline{320.1}$ ,  $\underline{435}/\underline{325}$ ,  $\underline{435}/\underline{69.1}$ ,  $\underline{530}/\underline{350}$ 

### ABSTRACT:

This invention provides an isolated nucleic acid encoding a mammalian galanin receptor, an isolated galanin receptor protein, vectors comprising isolated nucleic acid encoding a mammalian galanin receptor, cells comprising such vectors, antibodies directed to a mammalian galanin receptor, nucleic acid probes useful for detecting nucleic acid encoding a mammalian galanin receptor, antisense oligonucleotides complementary to unique sequences of nucleic acid encoding a mammalian galanin receptor, nonhuman transgenic animals which express DNA encoding a normal or a mutant mammalian galanin receptor, as well as methods of determining binding of compounds to mammalian galanin receptors.

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	16. Document ID	): US 2	0030082201	A 1						
птэ:	Entry 16 of 78				rile:	PGPB		M	av 1.	2003

May 1, 2003

PGPUB-DOCUMENT-NUMBER: 20030082201

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030082201 A1

TITLE: Multivalent synthetic vaccine for cancer

PUBLICATION-DATE: May 1, 2003

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Mukherjee, Rama	Ghaziabad		IN	
Rao, M.R.S.	Bangalore		IN	
Burman, Arnand C.	Ghaziabad		IN	
Thomas, Becky	Ghaziabad		IЙ	
Prasad, Sudhanand	Ghaziabad		IN	
Sengupta, Paromita	Ghaziabad		IN	

h e b b g ee e f ef e heh b Record List Display Page 11 of 52

US-CL-CURRENT:  $\underline{424/190.1}$ ;  $\underline{435/252.33}$ ,  $\underline{435/320.1}$ ,  $\underline{435/6}$ ,  $\underline{435/69.3}$ ,  $\underline{530/350}$ ,  $\underline{536/23.2}$ 

#### ABSTRACT:

Multivalent vaccine comprising peptides from vasoactive intestinal peptide, bombesin, Substance P and epidermal growth factor are described. A method of constructing a multivalent gene for use in various expressions vectors and the protein recombinantly expressed in the prokaryotic expression systems are also described.

Full Title Citation Front Review Classification Date Reference Sequences Attachments 6000 Draw Desc

17. Document ID: US 20030050446 A1

L13: Entry 17 of 78

File: PGPB

Mar 13, 2003

PGPUB-DOCUMENT-NUMBER: 20030050446

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030050446 A1

TITLE: Regulation of human neuropeptide y-like g protein-coupled receptor

PUBLICATION-DATE: March 13, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Ramakrishnan, Shyam Brighton MA US

US-CL-CURRENT: <u>530/350</u>

### ABSTRACT:

Reagents which regulate human neuropeptide Y G protein-coupled receptor (NPY-GPCR) protein and reagents which bind to human NPY-GPCR gene products can play a role in preventing, ameliorating, or correcting dysfunctions or diseases including, but not limited to, obesity, diabetes, anxiety, hypertension, cocaine withdrawal, congestive heart failure, memory enhancement, cardiac and cerebral vasospasm, pheochromocytoma, ganglioneuroblastoma, Huntington's disease, Alzheimer' disease, and Parkinson's disease.

Fuil	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	<b>,</b> ,	KOMC	Drawn Desc

18. Document ID: US 20030049794 A1

L13: Entry 18 of 78

File: PGPB

Mar 13, 2003

PGPUB-DOCUMENT-NUMBER: 20030049794

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030049794 A1

TITLE: DNA encoding a human dopamine D1 receptor and uses thereof

PUBLICATION-DATE: March 13, 2003

INVENTOR-INFORMATION:

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RULE-47

NAME CITY STATE COUNTRY
Weinshank, Richard L. New York NY US
Hartig, Paul R. Kinnelon NJ US

US-CL-CURRENT: 435/69.1; 435/252.3, 435/254.2, 435/320.1, 435/325, 530/350, 536/23.5

#### ABSTRACT:

This invention provides isolated nucleic acid molecules encoding a human dopamine D.sub.1 receptor, isolated proteins which are human dopamine D.sub.1 receptor, vectors comprising isolated nucleic acid molecules encoding a human dopamine D.sub.1 receptor, mammalian cells comprising such vectors, antibodies directed to a human dopamine D.sub.1 receptor, nucleic acid probes useful for detecting nucleic acid encoding human dopamine D.sub.1 receptor, antisense oligonucleotides complementary to any sequences of a nucleic acid molecule which encodes a human dopamine D.sub.1 receptor, pharmaceutical compounds related to human dopamine D.sub.1 receptor, and nonhuman transgenic animals which express DNA a normal or a mutant human dopamine D.sub.1 receptor. This invention further provides methods for determining ligand binding, detecting expression, drug screening, and treatment involving a human dopamine D.sub.1 receptor.

Full	Title   Cit	ation Frent	Review	Classification	Date Refere	nos Sequences	: Attachments		KWIC	Draw Desi
	19. Do	cument ID	: US 20	0030027254	• • A1		-	**************		***************************************
L13:	Entry 1	9 of 78			Fi	le: PGPB			Feb 6,	2003

PGPUB-DOCUMENT-NUMBER: 20030027254

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030027254 A1

TITLE: Processes for preparing compositions involving GALR3 receptor specific

compounds

PUBLICATION-DATE: February 6, 2003

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Bard, Jonathan A.	Doylestown	PA	US	
Borowsky, Beth	Montclair	NJ	US	
Smith, Kelli E.	Fair Lawn	NJ	US	
Branchek, Theresa A.	Teaneck	NJ	US	
Gerald, Christophe P.G.	Ridgewood	NJ	US	•
Jones, Kenneth A.	Waltham	MA	US	

US-CL-CURRENT: <u>435/69.1</u>; <u>435/320.1</u>, <u>435/325</u>, <u>530/350</u>, <u>536/23.5</u>

### ABSTRACT:

This invention provides an isolated nucleic acid encoding a mammalian galanin receptor, an isolated galanin receptor protein, vectors comprising isolated nucleic acid encoding a mammalian galanin receptor, cells comprising such vectors, antibodies directed to a mammalian galanin receptor, nucleic acid probes useful for detecting nucleic acid encoding a mammalian galanin receptor, antisense oligonucleotides complementary to unique sequences of nucleic acid encoding a mammalian galanin

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receptor, nonhuman transgenic animals which express DNA encoding a normal or a mutant mammalian galanin receptor, as well as methods of determining binding of compounds to mammalian galanin receptors.

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | ----- KMC | Draw Des

20. Document ID: US 20030022277 A1

L13: Entry 20 of 78 File: PGPB Jan 30, 2003

PGPUB-DOCUMENT-NUMBER: 20030022277

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030022277 A1

TITLE: HUMAN NEUROPEPTIDE RECEPTOR

PUBLICATION-DATE: January 30, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

SOPPET, DANIEL R. CENTREVILLE VA US
LI, YI SUNNYVALE CA • US
ROSEN, CRAIG A. LAYTONSVILLE MD US

US-CL-CURRENT:  $\underline{435/69.1}$ ;  $\underline{435/320.1}$ ,  $\underline{435/325}$ ,  $\underline{435/6}$ ,  $\underline{435/7.1}$ ,  $\underline{530/350}$ ,  $\underline{536/23.5}$ 

### ABSTRACT:

The present invention relates to a novel human protein called human <u>neuropeptide</u> <u>receptor</u>, and isolated polynucleotides encoding this protein. Also provided are vectors, host cells, antibodies, and recombinant methods for producing this human protein. The invention further relates to diagnostic and therapeutic methods useful for diagnosing and treating disorders related to this novel human protein.

Full	Title	Citation	Fiont	Review	Classification	Date	Reference	Sequences	Attachmenta	,,	KWIC	Draw, Desc

21. Document ID: US 20030018184 A1

'L13: Entry 21 of 78 File: PGPB Jan 23, 2003

PGPUB-DOCUMENT-NUMBER: 20030018184

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030018184 A1

TITLE: Recombinant C140 receptor, its agonists and antagonists, and nucleic acids

encoding the receptor

PUBLICATION-DATE: January 23, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Sundelin, Johan Furulund CA SE Scarborough, Robert M. Belmont US

h eb b g e e e f e h eh ef b e

US-CL-CURRENT:  $\underline{536}/\underline{23.5}$ ;  $\underline{435}/\underline{320.1}$ ,  $\underline{435}/\underline{325}$ ,  $\underline{435}/\underline{455}$ ,  $\underline{435}/\underline{69.1}$ ,  $\underline{530}/\underline{350}$ 

#### ABSTRACT:

Nucleic acid molecules encoding the C140 cell surface receptor have been cloned and sequenced. The availability of C140 receptor DNA permits the recombinant production of the C140 receptor which can be produced on the surface of a cell, including an oocyte. The nucleic acid molecules are useful in an assay for detecting a substance which affects C140 receptor activity, either receptor agonists or antagonists. Further, the elucidation of the structure of the C140 receptor permits the design of agonist and antagonist compounds which are useful in such assays. The availability of the C140 receptor also permits production of antibodies specifically immunoreactive with one or more antigenic epitopes of the C140 receptor.

Full	Title Citation Fro	nt Review	Classification	Date -	Reference	Sequences	Attachments	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	KWIC	Draw Desc
	22. Document	: ID: US 20	0020182648	3 A1		***************************************		***************************************		***************************************
L13:	Entry 22 of 7	8			File:	PGPB		De	ec 5,	2002

PGPUB-DOCUMENT-NUMBER: 20020182648

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020182648 A1

TITLE: Ordered two-and three-dimensional structures of amphiphilic molecules

PUBLICATION-DATE: December 5, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Mojtabai, Fatemeh Demarest NJ US

US-CL-CURRENT: 435/7.9; 435/194, 435/287.2, 530/350

### ABSTRACT:

The invention pertains, at least in part, to a method for forming an ordered structure of amphiphilic molecules, such as proteins. The method includes contacting a population of amphiphilic molecules with a interface; compressing said population laterally to an appropriate pressure, such that an ordered structure at the interface is formed. The invention also pertains to the two- and three-dimensional ordered structures that are formed using the planar membrane compression method of the invention.

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23. Document ID: US 20020172940 A1

L13: Entry 23 of 78 File: PGPB Nov 21, 2002

PGPUB-DOCUMENT-NUMBER: 20020172940

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020172940 A1

h eb b g ee ef e h eh ef b e

TITLE: Methods and reagents for isolating biologically active peptides

PUBLICATION-DATE: November 21, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Gyuris, Jeno Winchester MA US Morris, Aaron J. Boston MA US

US-CL-CURRENT: 435/5; 435/7.1, 435/7.32, 436/518, 530/324, 530/350

### ABSTRACT:

One aspect of the present invention is the synthesis of a binary method that combines variegated peptide display libraries, e.g., in a "display mode", with soluble secreted peptide libraries, e.g., in a "secretion mode", to yield a method for the efficient isolation of peptides having a desired biological activity.

1	Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	 KNMC   Drawn Desc
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# 24. Document ID: US 20020157119 A1

L13: Entry 24 of 78 File: PGPB Oct 24, 2002

PGPUB-DOCUMENT-NUMBER: 20020157119

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020157119 A1

TITLE: Identification of activated receptors and ion channels

PUBLICATION-DATE: October 24, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Beachy, Philip A. Towson MD US Taipale, Jussi Baltimore MD US

US-CL-CURRENT: 800/8; 435/194, 435/320.1, 435/354, 435/6, 435/7.1, 530/350

## ABSTRACT:

The present invention related to methods and reagents for generating and using activating mutations of receptors and ion channels.

Full   Title	e Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	 KMC	Draw Des

## 25. Document ID: US 20020150973 A1

L13: Entry 25 of 78 File: PGPB Oct 17, 2002

PGPUB-DOCUMENT-NUMBER: 20020150973

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020150973 A1

TITLE: Compositions and methods for the diagnosis and treatment of body weight disorders, including obesity

PUBLICATION-DATE: October 17, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Moore, Karen Maynard MA US Nagle, Deborah Lynn Watertown MA US

US-CL-CURRENT:  $\underline{435}/\underline{69.1}$ ;  $\underline{435}/\underline{183}$ ,  $\underline{435}/\underline{320.1}$ ,  $\underline{435}/\underline{325}$ ,  $\underline{435}/\underline{6}$ ,  $\underline{530}/\underline{350}$ ,  $\underline{536}/\underline{23.2}$ 

#### ABSTRACT:

The present invention relates to mammalian mahogany genes, including the human mahogany gene, which are novel genes involved in the control of mammalian body weight. The invention encompasses nucleotide sequences of the mahogany gene, host cell expression systems of the mahogany gene, and hosts which have been transformed by these expression systems, including transgenic animals. The invention also encompasses novel mahogany gene products, including mahogany proteins, polypeptides and peptides containing amino acid sequences mahogany proteins, fusion proteins of mahogany proteins polypeptides and peptides, and antibodies directed against such mahogany gene products. The present invention also relates to methods and compositions for the diagnosis and treatment of mammalian body weight disorders, including obesity, cachexia, and anorexia, and for the identification of subjects susceptible to such disorders. Further, the invention relates to methods of using the mahogany gene and gene products of the invention for the identification of compounds which modulate the expression of the mahogany gene and/or the activity of the mahogany gene product. Such compounds can be useful as therapeutic agents in the treatment of mammalian body weight disorders, including obesity, cachexia, and anorexia.

F	ull (	Title	Citation	Front	Review	Classification	Date	Reference	Attachments	 KWIC D	raw. Desc

26. Document ID: US 20020115155 A1

L13: Entry 26 of 78 File: PGPB Aug 22, 2002

PGPUB-DOCUMENT-NUMBER: 20020115155

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020115155 A1

TITLE: Human neuropeptide receptor

PUBLICATION-DATE: August 22, 2002

INVENTOR-INFORMATION:

NAME. CITY STATE COUNTRY RULE-47 Soppet, Daniel R. Centreville VΑ US Li, Yi Sunnyvale CA US Rosen, Craig A. Laytonsville MD US

US-CL-CURRENT: 435/69.1; 435/320.1, 435/325, 530/350, 536/23.2

ABSTRACT:

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Human <u>neuropeptide receptor</u> polypeptides and DNA (RNA) encoding such polypeptides and a procedure for producing such polypeptides by recombinant techniques is disclosed. Also disclosed are methods for utilizing such polypeptides for identifying antagonists and agonists to such polypeptides and methods of using the agonists and antagonists therapeutically to treat conditions related to the underexpression and overexpression of the <u>neuropeptide receptor</u> polypeptides, respectively. Also disclosed are diagnostic methods for detecting a mutation in the <u>neuropeptide receptor</u> nucleic acid sequences and an altered level of the soluble form of the receptors.

Full	Title Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	t*	KMC	Drawt Desc
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	27. Docum	nent ID:	US 2	0020115149	) A1						
L13:	Entry 27 of	f 78				File:	PGPB		Αu	ıa 22.	2002

PGPUB-DOCUMENT-NUMBER: 20020115149

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020115149 A1

TITLE: DNA encoding human 5-HT1D receptors and uses thereof

PUBLICATION-DATE: August 22, 2002

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Weinshank, Richard L.	New York	NY	US	
Branchek, Theresa	Teaneck	NJ	US	
Hartig, Paul R.	Mahwah	NJ	US	

US-CL-CURRENT: 435/69.1; 435/252.3, 435/320.1, 435/325, 530/350, 536/23.2

### ABSTRACT:

This invention provides isolated nucleic acid molecules encoding human 5-HT.sub.1D receptors, isolated proteins which are human 5-HT.sub.1D receptors, vectors comprising isolated nucleic acid molecules encoding human 5-HT.sub.1D receptors, mammalian cells comprising such vectors, antibodies directed to the human 5-HT.sub.1D receptors, nucleic acid probes useful for detecting nucleic acid encoding human 5-HT.sub.1D receptors, antisense oligonucleotides complementary to any sequences of a nucleic acid molecule which encodes a human 5-HT.sub.1D receptor, pharmaceutical compounds related to human 5-HT.sub.1D receptors, and nonhuman transgenic animals which express DNA a normal or a mutant human 5-HT.sub.1D receptor. This invention further provides methods for determining ligand binding, detecting expression, drug screening, and treatment involving the human 5-HT.sub.1D receptor.

Full Title	Citation Front Review	Classification   Date	Reference	Sequences	Attachments	hund	KOMC	Draw Desi
FF 28	Document ID: US 2					~~~~		······

20. Document ID. 05 20020070570 A

L13: Entry 28 of 78

File: PGPB

Jul 25, 2002

PGPUB-DOCUMENT-NUMBER: 20020098548

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020098548 A1

h eb bgeeef eheh ef be

TITLE: DNA encoding a human serotonin (5-HT2) receptor and uses thereof

PUBLICATION-DATE: July 25, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Kao, Hung-TehHackensackNJUSHartig, Paul R.MahwahNJUSBranchek, TheresaTeaneckNJUS

US-CL-CURRENT:  $\underline{435/69.1}$ ;  $\underline{435/235.1}$ ,  $\underline{435/320.1}$ ,  $\underline{435/325}$ ,  $\underline{530/350}$ ,  $\underline{536/23.5}$ 

#### ABSTRACT:

The present invention provides an isolated nucleic acid molecule encoding an 5-HT.sub.2 receptor, and an isolated protein which is a human 5-HT.sub.2 receptor.

The invention also provides vectors comprising DNA molecules encoding a human 5-HT.sub.2 receptor, and vectors adapted for expression of the 5-HT.sub.2 receptor in bacterial, yeast, or mammalian cells.

In addition, the invention provides a DNA probe useful for detecting nucleic acid encoding the 5-HT.sub.2 receptor, a method for determining whether a ligand which is not known to be capable of binding to the 5-HT.sub.2 receptor can bind to the 5-HT.sub.2 receptor, a method for detecting the presence of 5-HT.sub.2 receptor on the surface of a cell, and a method of screening drugs to identify drugs which specifically interact with, and bind to, the 5-HT.sub.2 receptor.

The invention herein also concerns an antibody directed to the human 5-HT.sub.2 receptor, such as a monoclonal antibody directed to an epitope of the 5-HT.sub.2 receptor present on the surface of a cell and having an amino acid sequence included within the amino acid sequence shown in FIG. 2.

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	 KMIC Draw Desc
										A - A - A

29. Document ID: US 20020094334 A1

L13: Entry 29 of 78 File: PGPB Jul 18, 2002

PGPUB-DOCUMENT-NUMBER: 20020094334

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020094334 A1

TITLE: Selective destruction of cells infected with human immunodeficiency virus

PUBLICATION-DATE: July 18, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Keener, William K. Idaho Falls ID US Ward, Thomas E. Idaho Falls ID US

US-CL-CURRENT: 424/160.1; 530/350, 530/359, 530/826

ABSTRACT:

h eb b g e e e f e h eh ef b e

Compositions and methods for selectively killing a cell containing a viral protease are disclosed. The composition is a variant of a protein synthesis inactivating toxin wherein a viral protease cleavage site is interposed between the A and B chains. The variant of the type II ribosome-inactivating protein is activated by digestion of the viral protease cleavage site by the specific viral protease. The activated ribosome-inactivating protein then kills the cell by inactivating cellular ribosomes. A preferred embodiment of the invention is specific for human immunodeficiency virus (HIV) and uses ricin as the ribosome-inactivating protein. In another preferred embodiment of the invention, the variant of the ribosome-inactivating protein is modified by attachment of one or more hydrophobic agents. The hydrophobic agent facilitates entry of the variant of the ribosome-inactivating protein into cells and can lead to incorporation of the ribosome-inactivating protein into viral particles. Still another preferred embodiment of the invention includes a targeting moiety attached to the variants of the ribosome-inactivating protein to target the agent to HIV infectable cells.

Full Title Citation Front	Review   Classification   Date   Reference	Sequences   /	Attachments	KWC Draw Des
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30. Document ID: US 20020082202 A1

L13: Entry 30 of 78 File: PGPB Jun 27, 2002

PGPUB-DOCUMENT-NUMBER: 20020082202

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020082202 A1

TITLE: Screening methods using ligands of the neutropeptide receptor HFGAN72

PUBLICATION-DATE: June 27, 2002

## INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Bergsma, Derk J.	Berwyn	PA	US	
Brooks, David P.	West Chester -	PA	US	
Gellai, Miklos	Devon	PA	US	
Wilson, Shelagh	Beckets Bramfield	TX	GB	-
Yanagisawa, Masashi	Dallas		US	

US-CL-CURRENT: <u>514/12</u>; <u>435/69.1</u>, <u>435/7.1</u>, <u>530/350</u>

## ABSTRACT:

Polypeptides of HFGAN72 receptor ligands and polynucleotides encoding the polypeptides are provided. Methods of using these polypeptides to diagnose diseases relating to the under- or over-expression of HFGAN72 receptor ligands are also provided. In addition, methods of identifying agonists or antagonists of the interaction of HFGAN72 receptor ligands with the HFGAN72 receptor are provided. Methods of treatment by administering the identified agonists or antagonists to patients in need thereof are further disclosed.

Full Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	 KAMC	Draw, Desi

11. Document ID: US 20020076755 A1

L13: Entry 31 of 78 File: PGPB Jun 20, 2002

PGPUB-DOCUMENT-NUMBER: 20020076755

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020076755 A1

TITLE: G protein coupled receptor (GPCR) agonists and antagonists and methods of

activating and inhibiting GPCR using the same

PUBLICATION-DATE: June 20, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Kuliopulos, Athan Winchester MA US Covic, Lidija Boston MA US

US-CL-CURRENT: <u>435/69.1</u>; <u>435/320.1</u>, <u>435/325</u>, <u>435/7.1</u>, <u>514/12</u>, <u>530/350</u>

#### ABSTRACT:

The invention relates generally to G protein coupled receptors and in particular to agonists and antagonists of G protein receptors and methods of using the same.

Full	Title Citation I	front Re	view Cl	lassification	Date	Reference	Sequences	Attachments	<b></b>	KWC	Draint Desc
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32. Document ID: US 20020061522 A1

L13: Entry 32 of 78 File: PGPB May 23, 2002

PGPUB-DOCUMENT-NUMBER: 20020061522

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020061522 A1

TITLE: 1983, 52881, 2398, 45449, 50289, and 52872 novel G protein-coupled receptors

and uses therefor

PUBLICATION-DATE: May 23, 2002

## INVENTOR-INFORMATION:

CITY NAME STATE COUNTRY RULE-47 Glucksmann, Maria Alexandra Lexington MΑ US Galvin, Katherine M. Jamaica Plain MA US Silos-Santiago, Inmaculada Cambridge ΜA US

US-CL-CURRENT: 435/6; 435/320.1, 435/325, 435/7.1, 530/350, 536/23.5

# ABSTRACT:

The invention provides isolated nucleic acids molecules, designated 1983, 52881, 2398, 45449, 50289, and 52872 nucleic acid molecules, which encode novel G protein-coupled receptor members. The invention also provides antisense nucleic acid molecules, recombinant expression vectors containing 1983, 52881, 2398, 45449, 50289, or 52872 nucleic acid molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 1983, 52881, 2398, 45449, 50289, or 52872 gene has been introduced or disrupted. The invention still further

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Record List Display Page 21 of 52

provides isolated 1983, 52881, 2398, 45449, 50289, or 52872 proteins, fusion proteins, antigenic peptides and anti-1983, 52881, 2398, 45449, 50289, or 52872 antibodies. Diagnostic methods utilizing compositions of the invention are also provided.

Full Title Citation Front Review Classification Date Reference Sequences Attachments ----

33. Document ID: US 20020056151 A1

L13: Entry 33 of 78 File: PGPB May 9, 2002

KWC Draw Desc

PGPUB-DOCUMENT-NUMBER: 20020056151

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020056151 A1

TITLE: Receptors for peptides from insects

PUBLICATION-DATE: May 9, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Antonicek, Horst-Peter Bergisch Gladbach DE Friedrich, Gabi Leverkusen DE Schulte, Thomas Koln DE

US-CL-CURRENT: 800/279; 435/320.1, 435/410, 435/69.1, 530/350, 536/23.5

ABSTRACT:

The invention relates to polypeptides having the biological activity of peptide receptors, and to nucleic acids encoding these polypeptides, and in particular to . their use for finding active compounds for crop protection.

Full Title Citation Front Review Classification Date Reference Sequences Attachments	Tit	le	Citati	n	Fiont	Revie	tun	Classification	Date	Reference	Sequences	Attachments	,y	٠٠	KMIC	Drawi Des
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34. Document ID: US 20020048791 A1

L13: Entry 34 of 78 File: PGPB Apr 25, 2002

PGPUB-DOCUMENT-NUMBER: 20020048791

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020048791 A1

TITLE: Human neuropeptide Y-like G protein-coupled receptor

PUBLICATION-DATE: April 25, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Zhelnin, Leonid Madison CT US Bloomquist, Brian T. New Haven CT US

US-CL-CURRENT:  $\underline{435}/\underline{69.1}$ ;  $\underline{435}/\underline{320.1}$ ,  $\underline{435}/\underline{325}$ ,  $\underline{435}/\underline{69.7}$ ,  $\underline{530}/\underline{350}$ ,  $\underline{536}/\underline{23.5}$ 

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## ABSTRACT:

Reagents which regulate human neuropeptide Y-like G protein-coupled receptor (NPY-like GPCR) protein and reagents which bind to human NPY-like GPCR gene products can play a role in preventing, ameliorating, or correcting dysfunctions or diseases including, but not limited to, obesity, diabetes, anxiety, hypertension, cocaine withdrawal, congestive heart failure, memory enhancement, cardiac and cerebral vasospasm, pheochromocytoma, ganglioneuroblastoma, Huntington's disease, Alzheimer's disease, and Parkinson's disease.

Full Title Citation Front Review Classification Date Reference Sequences Attachments - KWIC Draw Desc

35. Document ID: US 6800729 B2

L13: Entry 35 of 78

File: USPT

Oct 5, 2004

US-PAT-NO: 6800729

DOCUMENT-IDENTIFIER: US 6800729 B2

TITLE: Human G-Protein chemokine receptor HDGNR10 (CCR5 receptor)

DATE-ISSUED: October 5, 2004

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Li; Yi Gaithersburg MD Ruben; Steven M. Olney MD

US-CL-CURRENT: 530/350; 530/300

### ABSTRACT:

Human G-protein chemokine receptor polypeptides and DNA (RNA) encoding such polypeptides and a procedure for producing such polypeptides by recombinant techniques is disclosed. Also disclosed are methods for utilizing such polypeptides for identifying antagonists and agonists to such polypeptides and methods of using the agonists and antagonists therapeutically to treat conditions related to the underexpression and overexpression of the G-protein chemokine receptor polypeptides, respectively. Also disclosed are diagnostic methods for detecting a mutation in the G-protein chemokine receptor nucleic acid sequences and detecting a level of the soluble form of the receptors in a sample derived from a host.

59 Claims, 4 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 4

Full Title Citation Front Review Classification Date Reference KindC Draw. Desc

36. Document ID: US 6770449 B2

L13: Entry 36 of 78 File: USPT Aug 3, 2004

US-PAT-NO: 6770449

DOCUMENT-IDENTIFIER: US 6770449 B2

Record List Display Page 23 of 52

TITLE: Methods of assaying receptor activity and constructs useful in such methods

DATE-ISSUED: August 3, 2004

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Barak; Lawrence S. Durham NC Caron; Marc G. Hillsborough NC

Ferguson; Stephen S. London CA

Zhang; Jie Durham NC

US-CL-CURRENT: 435/7.2; 435/325, 435/4, 435/7.1, 530/350

#### **ABSTRACT:**

Described are methods of detecting G-protein coupled receptor (GPCR) activity in vivo and in vitro; methods of assaying GPCR activity; and methods of screening for GPCR ligands, G protein-coupled receptor kinase (GRK) activity, and compounds that interact with components of the GPCR regulatory process. Constructs useful in such methods are described.

2 Claims, 21 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 14

Full Title Citation	Fiont	Review	Classification	Date Reference	,y	KMIC	Draw Des

37. Document ID: US 6750026 B2

L13: Entry 37 of 78 File: USPT Jun 15, 2004

US-PAT-NO: 6750026

DOCUMENT-IDENTIFIER: US 6750026 B2

TITLE: Screening methods using ligands of the neutropeptide receptor HFGAN72

DATE-ISSUED: June 15, 2004

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY
Bergsma; Derk J. Berwyn PA
Brooks; David P. West Chester PA
Gellai; Miklos Devon PA

Wilson; Shelagh Beckets Bramfield GB

Yanagisawa; Masashi Dallas TX

US-CL-CURRENT:  $\underline{435/7.1}$ ;  $\underline{435/252.3}$ ,  $\underline{435/325}$ ,  $\underline{435/4}$ ,  $\underline{530/300}$ ,  $\underline{530/324}$ ,  $\underline{530/399}$ 

### ABSTRACT:

Polypeptides of HFGAN72 receptor ligands and polynucleotides encoding the polypeptides are provided. Methods of using these polypeptides to diagnose diseases relating to the under- or over-expression of HFGAN72 receptor ligands are also provided. In addition, methods of identifying agonists or antagonists of the interaction of HFGAN72 receptor ligands with the HFGAN72 receptor are provided.

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Record List Display Page 24 of 52

Methods of treatment by administering the identified agonists or antagonists to patients in need thereof are further disclosed.

2 Claims, 5 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 5

Full Title Citation Front Review Classification Date Reference ______ KMC Draw Desc

38. Document ID: US 6743594 B1

L13: Entry 38 of 78

File: USPT

Jun 1, 2004

US-PAT-NO: 6743594

DOCUMENT-IDENTIFIER: US 6743594 B1

TITLE: Methods of screening using human G-protein chemokine receptor HDGNR10 (CCR5)

DATE-ISSUED: June 1, 2004

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Li; Yi Gaithersburg MD Ruben; Steven M. Olney MD

US-CL-CURRENT:  $\underline{435}/\underline{7.2}$ ;  $\underline{435}/\underline{252.3}$ ,  $\underline{435}/\underline{320.1}$ ,  $\underline{435}/\underline{69.1}$ ,  $\underline{530}/\underline{350}$ ,  $\underline{536}/\underline{23.1}$ 

### ABSTRACT:

Human G-protein chemokine receptor polypeptides and DNA (RNA) encoding such polypeptides and a procedure for producing such polypeptides by recombinant techniques is disclosed. Also disclosed are methods for utilizing such polypeptides for identifying antagonists and agonists to such polypeptides and methods of using the agonists and antagonists therapeutically to treat conditions related to the underexpression and overexpression of the G-protein chemokine receptor polypeptides, respectively. Also disclosed are diagnostic methods for detecting a mutation in the G-protein chemokine receptor nucleic acid sequences and detecting a level of the soluble form of the receptors in a sample derived from a host.

66 Claims, 4 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 4

Full Title Citation Front Review Classification Date Reference KNIC Drain, Desc

39. Document ID: US 6737408 B1

L13: Entry 39 of 78

File: USPT

May 18, 2004

US-PAT-NO: 6737408

DOCUMENT-IDENTIFIER: US 6737408 B1

TITLE: Compounds for control of appetite, blood pressure, cardiovascular response,

libido, and circadian rhythm

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DATE-ISSUED: May 18, 2004

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Balasubramanium; Ambikaipakan Cincinnati OH Chance; William T. Withamsville OH

US-CL-CURRENT: 514/18; 435/7.1, 514/19, 530/300, 530/331, 530/335, 530/344, 530/345

### ABSTRACT:

This invention relates generally to dipeptides and tripeptides and to methods for pharmaceutical treatment of mammals using analogs of such dipeptides and tripeptides. More specifically, the invention relates to tripeptides and their analogs, to pharmaceutical compositions containing such dipeptides and tripeptides and to methods of treatment of mammals using such dipeptides and tripeptides. In addition, the invention relates to methods of treatment of mammals using such dipeptides and tripeptides for control of appetite, blood pressure, cardiovascular response, libido, and circadian rhythm.

12 Claims, 3 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 2

Full	Title   Citation   Front   I	Review   Classification	Date Reference		,,	KWC	Draw Desc
	40. Document ID:	US 6733990 B1					
L13:	Entry 40 of 78		File:	USPT	May	11,	2004

US-PAT-NO: 6733990

DOCUMENT-IDENTIFIER: US 6733990 B1

TITLE: Nucleic acid encoding 15571, a GPCR-like molecule of the secretin-like family

DATE-ISSUED: May 11, 2004

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Hodge; Martin R. Arlington MA

Lloyd; Clare London GB

Weich; Nadine S. Brookline MA

US-CL-CURRENT: 435/69.1; 435/252.3, 435/254.11, 435/320.1, 435/325, 435/471, 435/71.1, 435/71.2, 530/350, 536/23.5

## ABSTRACT:

Novel GPCR-like polypeptides, proteins, and nucleic acid molecules are disclosed. In addition to isolated, full-length GPCR-like proteins, the invention further provides isolated GPCR-like fusion proteins, antigenic peptides, and anti-GPCR-like antibodies. The invention also provides GPCR-like nucleic acid molecules, recombinant expression vectors containing a nucleic acid molecule of the invention, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a GPCR-like gene has been introduced or disrupted. Diagnostic, screening, and therapeutic methods utilizing compositions of the invention are also

provided.

12 Claims, 28 Drawing figures Exemplary Claim Number: 1
Number of Drawing Sheets: 28

Full Title Citation Front Review Classification Date Reference KWC Draw Desc

11. Document ID: US 6727348 B2

L13: Entry 41 of 78

File: USPT

Apr 27, 2004

US-PAT-NO: 6727348

DOCUMENT-IDENTIFIER: US 6727348 B2

TITLE: Compositions and methods for the diagnosis and treatment of body weight

disorders, including obesity

DATE-ISSUED: April 27, 2004

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Moore; Karen Maynard MA Nagle; Deborah Lynn Watertown MA

US-CL-CURRENT: <u>530/350</u>; <u>435/69.1</u>, 530/300, 536/23.1

### ABSTRACT:

The present invention relates to mammalian mahogany genes, including the human mahogany gene, which are novel genes involved in the control of mammalian body weight. The invention encompasses nucleotide sequences of the mahogany gene, host cell expression systems of the mahogany gene, and hosts which have been transformed by these expression systems, including transgenic animals. The invention also encompasses novel mahogany gene products, including mahogany proteins, polypeptides and peptides containing amino acid sequences mahogany proteins, fusion proteins of mahogany proteins polypeptides and peptides, and antibodies directed against such mahogany gene products. The present invention also relates to methods and compositions for the diagnosis and treatment of mammalian body weight disorders, including obesity, cachexia, and anorexia, and for the identification of subjects susceptible to such disorders. Further, the invention relates to methods of using the mahogany gene and gene products of the invention for the identification of compounds which modulate the expression of the mahogany gene and/or the activity of the mahogany gene product. Such compounds can be useful as therapeutic agents in the treatment of mammalian body weight disorders, including obesity, cachexia, and anorexia.

17 Claims, 181 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 173

Full	Title	Citation	Front	Review	Classification	Date	Reference	enna	KMMC Draw Desi

42. Document ID: US 6713277 B1

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L13: Entry 42 of 78

File: USPT

Mar 30, 2004

US-PAT-NO: 6713277

DOCUMENT-IDENTIFIER: US 6713277 B1

TITLE: Methods and composition for the diagnosis and treatment of body weight

disorders, including obesity

DATE-ISSUED: March 30, 2004

INVENTOR-INFORMATION:

NAME CITY

STATE ZIP CODE

COUNTRY

Moore; Karen

Maynard

MA

Nagle; Deborah Lynn

Watertown

MΑ

US-CL-CURRENT:  $\underline{435}/\underline{69.1}$ ;  $\underline{435}/\underline{252.3}$ ,  $\underline{435}/\underline{320.1}$ ,  $\underline{530}/\underline{350}$ ,  $\underline{536}/\underline{23.1}$ 

#### ABSTRACT:

The present invention relates to mammalian mahogany genes, including the human mahogany gene, which are novel genes involved in the control of mammalian body weight. The invention encompasses nucleotide sequences of the mahogany gene, host cell expression systems of the mahogany gene, and hosts which have been transformed by these expression systems, including transgenic animals. The invention also encompasses novel mahogany gene products, including mahogany proteins, polypeptides and peptides containing amino acid sequences mahogany proteins, fusion proteins of mahogany proteins polypeptides and peptides, and antibodies directed against such mahogany gene products. The present invention also relates to methods and compositions for the diagnosis and treatment of mammalian body weight disorders, including obesity, cachexia, and anorexia, and for the identification of subjects susceptible to such disorders. Further, the invention relates to methods of using the mahogany gene and gene products of the invention for the identification of compounds which modulate the expression of the mahogany gene and/or the activity of the mahogany gene product. Such compounds can be useful as therapeutic agents in the treatment of mammalian body weight disorders, including obesity, cachexia, and anorexia.

28 Claims, 183 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 173

Full	Title C	tation	Front	Review	Classification	Date	Reference		KWAC	
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## 43. Document ID: US 6641820 B1

L13: Entry 43 of 78

File: USPT

Nov 4, 2003

US-PAT-NO: 6641820

DOCUMENT-IDENTIFIER: US 6641820 B1

TITLE: Clostridial toxin derivatives and methods to treat pain

DATE-ISSUED: November 4, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

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Oct 21, 2003

Donovan; Stephen

Capistrano Beach

CA

US-CL-CURRENT: 424/239.1; 435/252.3, 435/320.1, 435/325, 435/69.1, 435/69.7, 435/70.1, 514/12, 514/14, 514/2, 530/350, 530/412

### ABSTRACT:

Methods for treating a bone tumor, in particular pain associated with bone tumor, by administration to a patient of a therapeutically effective amount of an agent are disclosed. The agent may include a clostridial neurotoxin component attached to a targeting moiety, wherein the targeting moiety is selected from the group consisting of transmission compounds which can be released from neurons upon the transmission of pain signals by the neurons, and compounds substantially similar to the transmission compounds.

8 Claims, 0 Drawing figures Exemplary Claim Number: 1

Full	Title	Citation Front Review	Classification	Date	Reference	,	KWIC	Draw Desc
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	44.	Document ID: US	5635432 B1					

File: USPT

US-PAT-NO: 6635432

L13: Entry 44 of 78

DOCUMENT-IDENTIFIER: US 6635432 B1

TITLE: Peptide potentiation of acid-sensory ion channel in pain

DATE-ISSUED: October 21, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Welsh; Michael J. Riverside IA
Askwith; Candice C. Iowa City IA

US-CL-CURRENT: 435/7.21; 435/252.3, 435/320.1, 435/4, 435/6, 435/69.1, 436/501, 530/300, 530/350, 536/23.5

### ABSTRACT:

An assay for determining agonists, antagonists, or modulators for acid-sensing ion channels. The assay is especially useful for screening analgesics. The screening assay can be provided in a kit form. The assay comprises administering the composition to be screened to cells expressing acid-gated channels and then determining whether the composition inhibits, enhances, or has no effect on the channels when acid is introduced. The determination can be performed by analyzing whether a current is sustained by the cells in the presence of the composition and the acid. This current can be compared to that sustained by the FMRFamide and related peptides.

11 Claims, 26 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 9 45. Document ID: US 6632621 B1

L13: Entry 45 of 78

File: USPT

Oct 14, 2003

US-PAT-NO: 6632621

DOCUMENT-IDENTIFIER: US 6632621 B1

** See image for <u>Certificate of Correction</u> **

TITLE: G protein-coupled receptor-like receptors and modulators thereof

DATE-ISSUED: October 14, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY Lowery; David E. Portage Geary; Timothy G. Kalamazoo ΜT Kubiak; Teresa M. Richland MI Larsen; Martha J. Kalamazoo. MI

US-CL-CURRENT:  $\underline{435}/\underline{7.22}$ ;  $\underline{435}/\underline{7.2}$ ,  $\underline{435}/\underline{7.21}$ ,  $\underline{530}/\underline{350}$ ,  $\underline{530}/\underline{388.6}$ ,  $\underline{536}/\underline{23.7}$ ,  $\underline{930}/\underline{210}$ 

### ABSTRACT:

The invention provides neuropeptide ligands, G protein-coupled receptors and methods of screening for modulators of receptor activity. Identified modulators, including neuropeptide ligand mimetics, are useful as biostatic and biocidal agents of varying scope, ranging from lethal activity restricted to particular invertebrate parasites to broad spectrum invertebrate parasiticides active on a wide range of invertebrates, including helminths and insects.

21 Claims, 1 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 1

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# 15 46. Document ID: US 6627197 B2

L13: Entry 46 of 78

File: USPT

Sep 30, 2003

US-PAT-NO: 6627197

DOCUMENT-IDENTIFIER: US 6627197 B2

TITLE: Selective destruction of cells infected with human immunodeficiency virus

DATE-ISSUED: September 30, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

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US-CL-CURRENT: 424/183.1; 424/94.1, 435/23, 514/2, 530/327, 530/350, 530/370, 530/377, 530/826

## ABSTRACT:

Compositions and methods for selectively killing a cell containing a viral protease are disclosed. The composition is a variant of a protein synthesis inactivating toxin wherein a viral protease cleavage site is interposed between the A and B chains. The variant of the type II ribosome-inactivating protein is activated by digestion of the viral protease cleavage site by the specific viral protease. The activated ribosome-inactivating protein then kills the cell by inactivating cellular ribosomes. A preferred embodiment of the invention is specific for human immunodeficiency virus (HIV) and uses ricin as the ribosome-inactivating protein. In another preferred embodiment of the invention, the variant of the ribosome-inactivating protein is modified by attachment of one or more hydrophobic agents. The hydrophobic agent facilitates entry of the variant of the ribosome-inactivating protein into cells and can lead to incorporation of the ribosome-inactivating protein into viral particles. Still another preferred embodiment of the invention includes a targeting moiety attached to the variants of the ribosome-inactivating protein to target the agent to HIV infectable cells.

43 Claims, 0 Drawing figures Exemplary Claim Number: 1

Full Title	Citation Front	Review	Classification	Date	Reference		,,	KMAC	Dram. Des
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L13: Entry 47 of 78

File: USPT

Aug 19, 2003

US-PAT-NO: 6608025

DOCUMENT-IDENTIFIER: US 6608025 B1

TITLE: Human NESP55 polypeptides, polynucleotides and uses thereof

DATE-ISSUED: August 19, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Fraser; Douglas Nottingham GB
St. Gallay; Steven Nottingham GB

US-CL-CURRENT: 514/2; 514/16, 514/18, 530/300, 530/328, 530/330, 530/350

### ABSTRACT:

A substantially pure polypeptide (human NESP55) comprising the amino acid sequence

(SEQ ID NO: 2) IRLEVPKRMDRRSRAQQWRRARHNYNDLCPPIGRRAATALLWLSCSIALL RALATSNARAQQRAAAQQRRSFLNAHHRSGAQVFPESPESESDHEHEEAD LELSLPECLEYEEEFDYETESEIESEIDFETEPETAPTTEPETEPE DDRGPVVPKHSTFGQSLTQRLHALKLRSPDASPSRAPPSTQEPQSPREGE ELKPEDKDPRRDPEESKEPKEEKQRRRCKPKKPTRRDASPESPSKKGPIP IRRH

or a variant, fragment, fusion or derivative thereof, or a fusion of a said variant or fragment or derivative, wherein the polypeptide variant has an amino acid sequence which has at least 90% identity with the amino acid sequence given above.

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Record List Display Page 31 of 52

NESP55 or fragments thereof may be useful in medicine for the treatment of obesity.

6 Claims, 2 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 2

Full Title Citation Front Review Classification Date Reference Miles

## 48. Document ID: US 6562945 B1

L13: Entry 48 of 78

File: USPT

May 13, 2003

US-PAT-NO: 6562945

DOCUMENT-IDENTIFIER: US 6562945 B1

TITLE: Galanin receptor

DATE-ISSUED: May 13, 2003

## INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY Shi-Hsiang; Shen Beaconsfield CA Sultan; Ahmad Dorval CA Wahlestedt; Claes Montreal CA Walker; Philippe Montreal CA

US-CL-CURRENT: 530/350; 435/320.1, 435/325, 435/69.1

## ABSTRACT:

The present invention is directed to a novel receptor for galanin which has been designated as galanin receptor 2. The invention encompasses both the receptor protein as well as nucleic acids encoding the protein. In addition, the present invention is directed to methods and compositions which rely upon either GAL-R2 proteins or nucleic acids.

8 Claims, 13 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 11

Full		Citation		Review	Classification	Date	Reference			41	KMC	Drawn Des
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## 49. Document ID: US 6555325 B1

L13: Entry 49 of 78

File: USPT

Apr 29, 2003

US-PAT-NO: 6555325

DOCUMENT-IDENTIFIER: US 6555325 B1

TITLE: System for detection of a functional interaction between a compound and a cellular signal transduction component

3

DATE-ISSUED: April 29, 2003

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INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Oehlen; Lambertus J. Tarrytown NY

US-CL-CURRENT: 435/7.31; 435/254.2, 435/350, 435/6, 435/69.1, 435/69.7, 435/7.1, 435/7.2, 530/350, 536/23.4, 536/23.5

#### ABSTRACT:

The present invention makes available a rapid, reproducible, robust assay system for screening and identifying pharmaceutically effective compounds that specifically interact with and modulate the activity of a cellular protein, e.g., a receptor or ion channel. The subject assay enables rapid screening of large numbers of compounds to identify those which act as an agonist or antagonist to the bioactivity of the cellular protein. In this system, the first cell is treated with a compound, and functional interaction of this compound with a cellular receptor yields a secreted signal. A second cell, bearing a receptor for this secreted signal, makes use of an indicator gene in a signaling pathway coupled to this second receptor. The subject assays include methods of identifying compounds which specifically modulate, for example, heterologous receptors coupled to the pheromone response pathway in yeast. The subject assays are particularly amenable to the identification of specific agonists and antagonists of G protein-coupled receptors.

11 Claims, 2 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 2

Full	Title   Citation   Front	Review CI	assification	Date	Reference			,,	KWIC	Draw Desc
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	50. Document II	D: US 650	4008 <b>B</b> 1				,			
L13:	Entry 50 of 78				File	USPT		Jan	17,	2003

US-PAT-NO: 6504008

DOCUMENT-IDENTIFIER: US 6504008 B1

TITLE: Cell based signal generation

DATE-ISSUED: January 7, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Xu; Jun Ossining NY
Trueheart; Joshua South Nyack NY

US-CL-CURRENT: <u>530</u>/<u>350</u>; <u>530</u>/<u>371</u>

## ABSTRACT:

The present invention makes available a rapid, reproducible, robust assay system for screening and identifying pharmaceutically effective compounds that specifically interact with and modulate the activity of a cellular protein, e.g., a receptor or ion channel. The subject assay enables rapid screening of large numbers of compounds to identify those which act as an agonist or antagonist to the bioactivity of the cellular protein. In particular, the assay of the invention makes use of a cell that harbors a protein that is responsive to a cellular signal transduction pathway. The protein is operatively linked to a polypeptide which causes a detectable signal to be

generated upon stimulation of the pathway, e.g., when a compound interacts with and modulates the activity of a cellular receptor or ion channel of the cell. Thus, the cell provides a signal generation means comprising a novel fusion protein the expression of which is independent of stimulation/activation of the signal transduction pathway, but the activity of which is responsive to the signal transduction pathway.

6 Claims, 0 Drawing figures Exemplary Claim Number: 1

Full Title Citation Front Review Classification Date Reference KMC Draw Desc

51. Document ID: US 6500436 B2

L13: Entry 51 of 78

File: USPT

Dec 31, 2002

US-PAT-NO: 6500436

DOCUMENT-IDENTIFIER: US 6500436 B2

TITLE: Clostridial toxin derivatives and methods for treating pain

DATE-ISSUED: December 31, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Donovan; Stephen

Capistrano Beach

CA

US-CL-CURRENT: 424/239.1; 435/252.3, 435/320.1, 435/325, 435/68.1, 435/69.1,

435/70.1, 514/12, 514/2, 530/350, 530/412, 536/23.1

### ABSTRACT:

Agents for treating pain, methods for producing the agents and methods for treating pain by administration to a patient of a therapeutically effective amount of the agent. The agent can include a clostridial neurotoxin, or a component or fragment or derivative thereof, attached to a targeting moiety, wherein the targeting moiety is selected from a group consisting of transmission compounds which can be released from neurons upon the transmission of pain signals by the neurons, and compounds substantially similar to the transmission compounds.

22 Claims, 0 Drawing figures Exemplary Claim Number: 1

Full Title Citation Front Review Classification Date Reference KWC Draw Des

52. Document ID: US 6468767 B1

L13: Entry 52 of 78

File: USPT

Oct 22, 2002

US-PAT-NO: 6468767

DOCUMENT-IDENTIFIER: US 6468767 B1

TITLE: DNA encoding a human dopamine D1 receptor and uses thereof

h e b b g e e e f e h eh ef b

DATE-ISSUED: October 22, 2002

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Weinshank; Richard L. New York NY Hartig; Paul R. Kinnelon NJ

US-CL-CURRENT: <u>435/69.1</u>; <u>530/350</u>

#### ABSTRACT:

This invention provides isolated nucleic acid molecules encoding a human dopamine D.sub.1 receptor, isolated proteins which are human dopamine D.sub.1 receptor, vectors comprising isolated nucleic acid molecules encoding a human dopamine D.sub.1 receptor, mammalian cells comprising such vectors, antibodies directed to a human dopamine D.sub.1 receptor, nucleic acid probes useful for detecting nucleic acid encoding human dopamine D.sub.1 receptor, antisense oligonucleotides complementary to any sequences of a nucleic acid molecule which encodes a human dopamine D.sub.1 receptor, pharmaceutical compounds related to human dopamine D.sub.1 receptor, and nonhuman transgenic animals which express DNA a normal or a mutant human dopamine D.sub.1 receptor. This invention further provides methods for determining ligand binding, detecting expression, drug screening, and treatment involving a human dopamine D.sub.1 receptor.

3 Claims, 10 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 10

Full	Title Citation	Front	Review	Classification	Date	Reference		<i>,</i> ,	KMC	Draw. D
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53. Document ID: US 6441133 B1

L13: Entry 53 of 78 File: USPT Aug 27, 2002

US-PAT-NO: 6441133

DOCUMENT-IDENTIFIER: US 6441133 B1

TITLE: Thyrotropin-releasing hormone receptor 2(TRHR-2)

DATE-ISSUED: August 27, 2002

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Walker; Philippe Montreal CA

US-CL-CURRENT: 530/350; 435/69.1

## ABSTRACT:

The present invention is directed to the novel receptor for TRH which has been designated as TRH receptor 2. The invention encompasses both the receptor protein as well as nucleic acids encoding the protein. In addition, the present invention is directed to methods and compositions which rely upon either TRHR-2 proteins or nucleic acids.

8 Claims, 0 Drawing figures

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Exemplary Claim Number: 1

Full Title Citation Front Review Classification Date Reference KMiC Draw, Desc

54. Document ID: US 6423504 B1

L13: Entry 54 of 78

File: USPT

ZIP CODE

Jul 23, 2002

US-PAT-NO: 6423504

DOCUMENT-IDENTIFIER: US 6423504 B1

TITLE: Human-derived bradeion proteins, DNA coding for the proteins, and uses thereof

DATE-ISSUED: July 23, 2002

INVENTOR-INFORMATION:

NAME CITY

STATE

COUNTRY

Tanaka; Manami Tanaka; Tomoo

Ibaraki Kanagawa JР JP , .

US-CL-CURRENT:  $\underline{435}/\underline{7.23}$ ;  $\underline{424}/\underline{138.1}$ ,  $\underline{435}/\underline{320.1}$ ,  $\underline{435}/\underline{6}$ ,  $\underline{435}/\underline{69.1}$ ,  $\underline{530}/\underline{350}$ ,  $\underline{530}/\underline{388.1}$ ,

<u>530/389.1</u>, <u>536/23.5</u>

### ABSTRACT:

A human-derived bradeion protein, which has the following properties: (i) it is a transmembranous protein; (ii) it has a structure characteristic of growth hormone and cytokine receptors even in a structure of its transmembranous portion when its structure is determined by a hydrophobicity analysis according to Kyte-Doolittle method; (iii) it is expressed at a high level in a human adult brain, and in less amount in the heart, while it is not expressed in other adult organs or fetus; (iv) it induces programmed cell death (apoptosis) when over-expressed in a cultured human nerve cell lines; (v) it induces termination of cell division and aging when overexpressed in a cultured human normal cell; (vi) it is located in cytoplasm, and forms an intracellular aggregate when overexpressed; and (vii) besides human adult neurons, it is specifically expressed in a human colorectal cancer cell line or in a skin cancer cell line, or an analogue thereof.

18 Claims, 8 Drawing figures Exemplary Claim Number: 1,13 Number of Drawing Sheets: 7

Full Title Citation Front	Review Classification	Date Reference	) <b>4</b>	KNMC Drawn Desi
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55. Document ID: US 6388055 B1

L13: Entry 55 of 78

File: USPT

May 14, 2002

US-PAT-NO: 6388055

DOCUMENT-IDENTIFIER: US 6388055 B1

TITLE: Mouse CC-CKR5 receptor polypeptide

DATE-ISSUED: May 14, 2002

e b b g ee e f e h eh ef INVENTOR-INFORMATION:

NAME

CITY

STATE Z

ZIP CODE

COUNTRY

Bergsma; Derk J.

Berwyn

PΑ

Brawner; Mary E.

Berwyn

PA

Shabon; Usman

Swarthmore

PA

US-CL-CURRENT: 530/350; 530/351

## ABSTRACT:

Mouse CC-CKR5 polypeptides and DNA (RNA) encoding such mouse CC-CKR5 and a procedure for producing such polypeptides by recombinant techniques is disclosed. Also disclosed are methods for utilizing such mouse CC-CKR5 in the development of gene knockout mice for use as a model for human immunodeficiency virus.

2 Claims, 2 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 3

Full	Title   Citatio	n Fior	nt   Review	Classification	Date	Reference		****** . ·	KWIC	Draw Desc
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56. Document ID: US 6383762 B1

L13: Entry 56 of 78

File: USPT

May 7, 2002

US-PAT-NO: 6383762

DOCUMENT-IDENTIFIER: US 6383762 B1

TITLE: Methods of obtaining compounds that interact with a human serotonin (5-HT2)

receptor

DATE-ISSUED: May 7, 2002

INVENTOR-INFORMATION:

NAME

_ _ .

Hackensack

STATE ZIP CODE

COUNTRY

Kao; Hung-Teh
Hartig; Paul R.

Mahwah

CITY

NJ NJ

Branchek; Theresa

Teaneck

NJ

US-CL-CURRENT: 435/7.21; 435/325, 435/69.1, 530/350, 536/23.5

### ABSTRACT:

The present invention provides a method of obtaining a composition which comprises determining whether a chemical compound binds to a human 5-HT.sub.2 receptor expressed on the surface of a mammalian cell transfected with a vector adapted for expressing the receptor in the cell, and if the compound binds to the receptor, admixing the compound with a carrier. The present invention further provides a method of obtaining a composition which comprises determining whether a chemical compound binds to and activates or binds to and inhibits activation of a human 5-HT.sub.2 receptor expressed on the surface of a mammalian cell, wherein the human 5-HT.sub.2 receptor is expressed on the surface of a mammalian cell transfected with a vector adapted for expressing the receptor in the cell, and if the compound binds to and activates or binds to and inhibits activation of the receptor, admixing the compound with a carrier.

5 Claims, 4 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 10

Full Title Citation Front Review Classification Date Reference

57. Document ID: US 6348574 B1

L13: Entry 57 of 78

File: USPT

Feb 19, 2002

US-PAT-NO: 6348574

DOCUMENT-IDENTIFIER: US 6348574 B1

TITLE: Seven transmembrane receptors

DATE-ISSUED: February 19, 2002

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Godiska; Ronald Bothell WA
Gray; Patrick W. Seattle WA
Schweickart; Vicki Louise Seattle WA

US-CL-CURRENT: 530/350; 530/388.22, 536/23.5

## ABSTRACT:

DNA sequences encoding seven novel seven transmembrane receptors and variants thereof are disclosed as well as materials and methods for production of the same by recombinant techniques. Antibody substances specific for each of the seven transmembrane receptors are disclosed as useful for the modulation of the ligand/receptor binding reactions of the receptors.

17 Claims, 2 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 2

Full Title	Citation Front	Classification	Reference	,	KWMC Drawa Des

58. Document ID: US 6258556 B1

L13: Entry 58 of 78

File: USPT

Jul 10, 2001

US-PAT-NO: 6258556

DOCUMENT-IDENTIFIER: US 6258556 B1

TITLE: cDNA and genomic clones encoding human .mu. opiate receptor and the purified

gene product

DATE-ISSUED: July 10, 2001

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

h e b b g e e e f e h eh ef b

IT

Uhl; George Towson MD Johnson; Peter Perry Hall MD

Persico; Antonio M. Milan

Wang; Jia Bei Baltimore MD

US-CL-CURRENT: 435/69.1; 435/320.1, 435/325, 530/350, 536/23.5, 536/24.31, 930/10

#### ABSTRACT:

A human .mu. opiate receptor cDNA has been identified from a cerebral cortical CDNA library using sequences from the rat .mu. opiate receptor CDNA. The human .mu. opiate receptor (h.mu.OR1) shares 95% amino acid identity with the rat sequence. The expressed .mu.OR1 recognizes tested opiate drugs and opioid peptides in a sodium— and GTP—sensitive fashion with affinities virtually identical to those displayed by the rat .mu. opiate receptor. Effects on cyclic AMP are similar to those noted for the rat .mu. opiate receptor. Overlapping genomic clones spanning 50 kilobasepairs and hybridizing with the h.mu.OR1 cDNA contains exon sequences encoding the entire open reading frame of the human A opiate receptor are described. Analysis of hybridization to DNA prepared from human rodent hybrid cell lines and chromosomal in situ hybridization studies indicate localization to 6q24-25. An MspI polymorphism, producing a 3.7 kb band, is being used to assess this gene's involvement in neuropsychiatric disorders involving opiatergic systems.

19 Claims, 6 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 3

Full Title Citation	Front Review	U Classification	Date	Reference	£1114	KARC Draw Desc
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### 59. Document ID: US 6255059 B1

L13: Entry 59 of 78

File: USPT

Jul 3, 2001

US-PAT-NO: 6255059

DOCUMENT-IDENTIFIER: US 6255059 B1

** See image for <u>Certificate of Correction</u> **

TITLE: Methods for identifying G protein coupled receptor effectors

DATE-ISSUED: July 3, 2001

### INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY Klein; Christine A. Ossining NY Murphy; Andrew J. M. Montclair NJ Fowlkes; Dana M. Chapel Hill NC Broach; James Princeton NJ Manfredi; John Ossining NY Paul; Jeremy Nyack NY Trueheart; Joshua South Nyack NY

US-CL-CURRENT:  $\underline{435}/\underline{7.31}$ ;  $\underline{435}/\underline{254.2}$ ,  $\underline{435}/\underline{254.21}$ ,  $\underline{435}/\underline{6}$ ,  $\underline{435}/\underline{69.1}$ ,  $\underline{435}/\underline{69.7}$ ,  $\underline{435}/\underline{7.2}$ ,  $\underline{530}/\underline{300}$ ,  $\underline{530}/\underline{350}$ ,  $\underline{536}/\underline{23.4}$ ,  $\underline{536}/\underline{23.5}$ 

## ABSTRACT:

heb bgeeef ehehefb

The present invention makes available a rapid, effective assay for screening and identifying pharmaceutically effective compounds that specifically interact with and modulate the activity of a cellular receptor or ion channel. The subject assay enables rapid screening of large numbers of polypeptides in a yeast expression library to identifying those polypeptides which induce or antagonize receptor bioactivity. The subject assay is particularly amenable for identifying surrogate ligands for orphan receptors.

18 Claims, 4 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 4

Full Title Citation Front Review Classification Date Reference 60. Document ID: US 6210967 B1.

US-PAT-NO: 6210967

L13: Entry 60 of 78

DOCUMENT-IDENTIFIER: US 6210967 B1

TITLE: DNA encoding a mammalian LPA receptor and uses thereof

DATE-ISSUED: April 3, 2001

INVENTOR-INFORMATION:

NAME

CITY

STATE

File: USPT

ZIP CODE

COUNTRY

Apr 3, 2001

Bard; Jonathan A.

Doylestown

PΑ

US-CL-CURRENT: 435/361; 435/252.3, 435/320.1, 435/325, 435/348, 435/356, 435/357, <u>435/365</u>, <u>435/366</u>, <u>530/350</u>, <u>536/23.5</u>, <u>536/24.31</u>

## ABSTRACT:

This invention provides an isolated nucleic acid encoding a mammalian LPA receptor, a purified mammalian LPA receptor, vectors comprising isolated nucleic acid encoding an mammalian LPA receptor, cells comprising such vectors, antibodies directed to a mammalian LPA receptor, nucleic acid probes useful for detecting nucleic acid encoding a mammalian LPA receptor, antisense oligonucleotides complementary to unique sequences of nucleic acid encoding mammalian LPA receptor, transgenic, nonhuman animals which express DNA encoding a normal or a mutant mammalian LPA receptor, methods of isolating an mammalian LPA receptor, methods of treating an abnormality that is linked to the activity of the mammalian LPA receptor, as well as methods of determining binding of compounds to mammalian LPA receptors.

21 Claims, 11 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 11

Full	Title Citation	Front	Review	Classification	Date	Reference	press	KWAC	Draw, Desc
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61. Document ID: US 6133420 A

L13: Entry 61 of 78

File: USPT

Oct 17, 2000

e b b g ee e f ef e heh b US-PAT-NO: 6133420

DOCUMENT-IDENTIFIER: US 6133420 A

TITLE: GPR14 polypeptides

DATE-ISSUED: October 17, 2000

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Ames, Jr.; Robert S. Havertown PA
Sarau; Henry M. Harleysville PA
Foley; James J. Radnor PA
Shabon; Usman Swarthmore PA
Bergsma; Derk Berwyn PA

Chambers; Jonathan K. Haslingfield GB

US-CL-CURRENT: 530/350; 435/69.1, 530/300, 530/324, 530/326

## ABSTRACT:

Human GPR14 polypeptides and polynucleotides and methods for producing such polypeptides by recombinant techniques are disclosed. Also disclosed are methods for utilizing Human GPR14 polypeptides and polynucleotides in the design of protocols for the treatment of infections such as bacterial, fungal, protozoan and viral infections, particularly infections caused by HIV-1 or HIV-2; pain; cancers; anorexia; bulimia; asthma; Parkinson's disease; acute heart failure; hypotension; hypertension; urinary retention; osteoporosis; angina pectoris; myocardial infarction; ulcers; asthma; allergies; benign prostatic hypertrophy; and psychotic and neurological disorders, including anxiety, schizophrenia, manic depression, delirium, dementia, severe mental retardation and dyskinesias, such as Huntington's disease or Gilles dela Tourett's syndrome, among others and diagnostic assays for such conditions.

7 Claims, 8 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 7

Full 7 Title 1	Citation	Front	Review	Classification	Date	Reference		,	KWIC	Drawi Des
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# 62. Document ID: US 6087115 A

L13: Entry 62 of 78

File: USPT

Jul 11, 2000

US-PAT-NO: 6087115

DOCUMENT-IDENTIFIER: US 6087115 A

TITLE: Methods of identifying negative antagonists for G protein coupled receptors

DATE-ISSUED: July 11, 2000

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Gershengorn; Marvin C. New York NY
Arvanitakis; Leandros New York NY
Geras-Raaka; Elizabeth Dobbs Ferry NY

h e b b g e e e f e h eh ef b e

Cesarman; Ethel

Hoboken

NJ

US-CL-CURRENT: <u>435/7.21</u>; <u>435/252.3</u>, <u>435/254.11</u>, <u>435/325</u>, <u>435/365</u>, <u>435/6</u>, <u>435/69.1</u>, <u>435/7.2</u>, <u>435/8</u>, <u>530/350</u>, <u>536/23.1</u>, <u>536/23.72</u>, <u>536/24.1</u>

#### ABSTRACT:

The present invention is directed to a constitutively active G protein coupled receptor of human herpesvirus 8, as well as a method of identifying negative antagonists of a constitutively active G protein coupled receptor. The method comprises co-expressing in a host cell a constitutively active G protein coupled receptor and a reporter protein, wherein expression of the reporter protein is controlled by a promoter responsive to a signalling pathway activated by the constitutively active G protein coupled receptor; exposing the host cell to a test substance; and determining a level of reporter protein activity, wherein the level of reporter protein activity indicates effectiveness of the test substance as a negative antagonist of the constitutively active G protein coupled receptor. The invention further provides a method of preventing tumor formation or cell proliferation caused by a constitutively active G protein coupled receptor. This method comprises administering an amount of the negative antagonist so identified to a subject in an amount effective to prevent tumor formation or cell proliferation.

7 Claims, 8 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 5

Full	Title	Citation	Front	Review	Classification	Date	Reference	,	 KWC Dram Desi
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63. Document ID: US 6005074 A

L13: Entry 63 of 78

File: USPT

Dec 21, 1999

US-PAT-NO: 6005074

DOCUMENT-IDENTIFIER: US 6005074 A

TITLE: Cloning of human GPR14 receptor

DATE-ISSUED: December 21, 1999

INVENTOR-INFORMATION:

NAME CITY

CITY

STATE

ZIP CODE

COUNTRY

Shabon; Usman

Swarthmore

PA

Bergsma; Derk

Berwyn

PΑ

US-CL-CURRENT: 530/350; 435/69.1, 530/324

### ABSTRACT:

Human GPR14 polypeptides and polynucleotides and methods for producing such polypeptides by recombinant techniques are disclosed. Also disclosed are methods for utilizing Human GPR14 polypeptides and polynucleotides in the design of protocols for the treatment of infections such as bacterial, fungal, protozoan and viral infections, particularly infections caused by HIV-1 or HIV-2; pain; cancers; anorexia; bulimia; asthma; Parkinson's disease; acute heart failure; hypotension; hypertension; urinary retention; osteoporosis; angina pectoris; myocardial infarction; ulcers; asthma; allergies; benign prostatic hypertrophy; and psychotic and neurological disorders; including anxiety, schizophrenia, manic depression,

Record List Display Page 42 of 52

delirium, dementia, severe mental retardation and dyskinesias, such as Huntington's disease or Gilles dela Tourett's syndrome, among others and diagnostic assays for such conditions.

4 Claims, 3 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 3

Full	Title	Citation	Fiont	Review	Classification	Date	Reference	,,	, KMC	Drawt Desc

# 64. Document ID: US 6001970 A

L13: Entry 64 of 78

File: USPT

Dec 14, 1999

US-PAT-NO: 6001970

DOCUMENT-IDENTIFIER: US 6001970 A

TITLE: Modified human neuropeptide Y1 Receptors

DATE-ISSUED: December 14, 1999

#### INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Cascieri; Margaret A. East Windsor NJ
MacNeil; Douglas John Westfield NJ
Strader; Catherine D. Verona NJ

US-CL-CURRENT: 530/350; 536/23.5

### ABSTRACT:

Modified neuropeptide Y receptors having deletions, replacements or additions in the third intracellular domain are identified and methods of making the modified receptors are provided. The invention includes the modified receptors, assays employing the modified receptors, cells expressing the modified receptors, compounds identified through the use of the modified receptors, including modulators of the receptors, and the use of the compounds to treat conditions, including obesity, diabetes, anxiety, hypertension, cocaine withdrawal, congestive heart failure, memory enhancement, cardiac and cerebral vasospasm, pheochromocytoma and ganglioneuroblastoma, and Huntington's, Alzheimer's and Parkinson's diseases.

2 Claims, 4 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 4



65. Document ID: US 6001963 A

L13: Entry 65 of 78

File: USPT

Dec 14, 1999

US-PAT-NO: 6001963

DOCUMENT-IDENTIFIER: US 6001963 A

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## Record List Display

TITLE: Ligands of the neuropeptide receptor HFGAN72

DATE-ISSUED: December 14, 1999

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Bergsma; Derk J. Berwyn PA Yanagisawa; Masashi Dallas TX

US-CL-CURRENT: 530/324; 530/300, 530/350, 530/399

#### ABSTRACT:

Polypeptides of HFGAN72 receptor ligands and polynucleotides encoding the polypeptides are provided. Methods of using these polypeptides to diagnose diseases relating to the under- or over-expression of HFGAN72 receptor ligands are also provided. In addition, methods of identifying agonists or antagonists of the interaction of HFGAN72 receptor ligands with the HFGAN72 receptor are provided. Methods of treatment by administering the identified agonists or antagonists to patients in need thereof are further disclosed.

3 Claims, 5 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 3

Full	Title Citation	Front	Review	Classification	Date	Reference		,,,,,,,	 KWC	Drawt Des
					ar .		 			

66. Document ID: US 5989545 A

L13: Entry 66 of 78 File: USPT Nov 23, 1999

US-PAT-NO: 5989545

DOCUMENT-IDENTIFIER: US 5989545 A

TITLE: Clostridial toxin derivatives able to modify peripheral sensory afferent

functions

DATE-ISSUED: November 23, 1999

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY
Foster; Keith Alan Wiltshire GB

Duggan; Michael John London GB

Shone; Clifford Charles Wiltshire GB

US-CL-CURRENT: 424/183.1; 424/832, 424/94.67, 435/220, 435/69.1, 435/69.7, 514/2, 530/350, 530/388.22, 530/391.7, 530/402

## ABSTRACT:

The invention relates to an agent specific for peripheral sensory afferents. The agent may inhibit the transmission of signals between a primary sensory afferent and a projection neuron by controlling the release of at least one neurotransmitter or neuromodulator from the primary sensory afferent. The agent may be used in or as a pharmaceutical for the treatment of pain, particularly chronic pain.

43 Claims, 4 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 4

Full Title Citation Front Review Classification Date Reference KWMC Draw Desc

67. Document ID: US 5976814 A

L13: Entry 67 of 78

File: USPT

Nov 2, 1999

US-PAT-NO: 5976814

DOCUMENT-IDENTIFIER: US 5976814 A

TITLE: DNA encoding a human neuropeptide Y/peptide YY/pancreatic polypeptide receptor

(Y4) and uses thereof

DATE-ISSUED: November 2, 1999

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Bard; Jonathan A. Wyckoff NJ Walker; Mary W. Elmwood Park ŊJ Branchek; Theresa Teaneck NJ Weinshank; Richard L. Teaneck ŊJ

US-CL-CURRENT: 435/7.2; 435/252.3, 435/320.1, 435/325, 435/69.1, 530/300, 530/350

## ABSTRACT:

This invention provides methods for determining whether a chemical compound specifically binds to and activates or inhibits activation of a human or rat Y4 receptor.

10 Claims, 9 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 19

Full Title Citation	Front Re	view Classification	Date Re			,,	KMC	Drawi Des
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☐ 68. Docur		JS 5958709 A	••••••	*************	 	**************		

US-PAT-NO: 5958709

DOCUMENT-IDENTIFIER: US 5958709 A

TITLE: Processes for identifying compounds that bind to the human Y4 receptor

DATE-ISSUED: September 28, 1999

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Bard; Jonathan A. Wyckoff

h e b g ee e f e heh ef b е Walker; Mary W.
Branchek; Theresa
Weinshank; Richard L.

Elmwood Park NJ
Teaneck NJ
New York NY

US-CL-CURRENT:  $\underline{435}/\underline{7.21}$ ;  $\underline{435}/\underline{325}$ ,  $\underline{435}/\underline{352}$ ,  $\underline{435}/\underline{363}$ ,  $\underline{435}/\underline{366}$ ,  $\underline{435}/\underline{69.1}$ ,  $\underline{530}/\underline{350}$ 

## ABSTRACT:

This invention provides an isolated nucleic acid molecule encoding a human Y4 receptor, an isolated protein which is a human Y4 receptor, vectors comprising an isolated nucleic acid molecule encoding a human Y4 receptors, mammalian cells comprising such vectors, antibodies directed to the human Y4 receptor, nucleic acid probes useful for detecting nucleic acid encoding human Y4 receptors, antisense oligonucleotides complementary to any sequences of a nucleic acid molecule which encodes a human Y4 receptor, pharmaceutical compounds related to human Y4 receptors, and nonhuman transgenic animals which express DNA a normal or a mutant human Y4 receptor. This invention further provides methods for determining ligand binding, detecting expression, drug screening, and treatment involving the human Y4 receptor.

6 Claims, 12 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 12

Full	Title	Citation	Front	Review	Classification	Date	Reference			, <u>,</u>	K	MC	Draw Desc
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	69.	Docum	ent ID	: US 5	932702 A			•					
L13:	Entr	y 69 of	78		•		File	: USPT			Aug	3,	1999

US-PAT-NO: 5932702

DOCUMENT-IDENTIFIER: US 5932702 A

** See image for <u>Certificate of Correction</u> **

TITLE: Human G-protein coupled receptor

DATE-ISSUED: August 3, 1999

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY Fujii; Ryo Ibaraki JP Hinuma; Shuji Ibaraki JP Li; Yi Gaithersburg MD Ruben; Steven M. Olney MD Soppet; Daniel R. Centreville VA

US-CL-CURRENT: 530/350; 435/69.1, 530/300, 530/324, 536/23.5

ABSTRACT:

Human G-protein coupled receptor polypeptides and DNA (RNA) encoding such polypeptides and a procedure for producing such polypeptides by recombinant techniques is disclosed. Also disclosed are methods for utilizing such polypeptides for identifying antagonists and agonists to such polypeptides and methods of using the agonists and antagonists therapeutically to treat conditions related to the underexpression and overexpression of the G-protein coupled receptor polypeptides, respectively. Also disclosed are diagnostic methods for detecting a mutation in the

G-protein coupled receptor nucleic acid sequences and an altered level of the soluble form of the receptors.

7 Claims, 3 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 3

Full	Title	Citation	Front	Review	Classification	Date	Reference	A	 KWIC Draw De

70. Document ID: US 5919901 A

L13: Entry 70 of 78

File: USPT

Jul 6, 1999

US-PAT-NO: 5919901

DOCUMENT-IDENTIFIER: US 5919901 A

TITLE: Neuropeptide Y receptor Y5 and nucleic acid sequences

DATE-ISSUED: July 6, 1999

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Hu; Yinghe	North Haven	CT		
McCaleb; Michael L.	Madison	CT		
Bloomquist; Brian T.	New Haven	CT		
Flores-Riveros; Jaime R.	Madison	CT		
Cornfield; Linda J.	Hamden	CT		

US-CL-CURRENT: 530/350; 530/395, 536/23.5

ABSTRACT:

The present invention provides novel NPY/PYY receptor proteins and the nucleic acid sequence encoding them. The invention is directed to the isolation, characterization, and pharmacological use of these receptors and nucleic acids. In particular, this invention provides human and rat NPY/PYY receptors (which we call the NPY Y5 receptor) and nucleic acids. Also provided are recombinant expression constructs useful for transfecting cells and expressing the protein in vitro and in vivo. The invention further provides methods for detecting expression levels of the protein as well as methods for screening for receptor antagonists and agonists to be used for the treatment of obesity or anorexia, respectively.

6 Claims, 2 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 2

Full	Title	Citation		Review	Classification	Date	Reference			·	KWAC	Dram Des
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71. Document ID: US 5888811 A

L13: Entry 71 of 78

File: USPT

Mar 30, 1999

US-PAT-NO: 5888811

DOCUMENT-IDENTIFIER: US 5888811 A

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Record List Display

TITLE: Corticotropin-releasing hormone receptor

DATE-ISSUED: March 30, 1999

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Largent; Brian Lee Chadds Ford PA
Chen; Ai-Ru Edison NJ
Kostich; Walter Alan Hockessin DE
Sperle; Karen Marie Hockessin DE

 $\text{US-CL-CURRENT: } \underline{435/320.1; } \underline{435/252.3}, \underline{435/254.11}, \underline{435/325}, \underline{435/69.1}, \underline{530/300}, \underline{530/350}, \\$

536/23.1, 536/23.5

ABSTRACT:

A novel human corticotropin releasing hormone (CRH) receptor which is a splice variant of the human CRH.sub.2 receptor subfamily and is designated human CRHR2.gamma. Fragments of CRHR2.gamma. Nucleic acid molecules which encode CRHR2.gamma and fragments, expression vectors comprising the nucleic acid molecules, and host cells containing the expression vectors. Antibodies and antibody fragments capable of binding the novel receptor. Nucleic acid molecules capable of hybridizing with the above nucleic acid molecules. Use of the novel receptor and receptor fragments, antibodies and antibody fragments in testing compounds for CRH antagonist activity and in treating diseases.

24 Claims, 0 Drawing figures Exemplary Claim Number: 13

Full	Title Citation	Front	Review	Classification	Date	Reference		*****	43	1000C	Drawt Des

72. Document ID: US 5885785 A

L13: Entry 72 of 78 File: USPT Mar 23, 1999

US-PAT-NO: 5885785

DOCUMENT-IDENTIFIER: US 5885785 A

TITLE: DNA encoding a human serotonin (5-HT2) receptor and uses thereof

DATE-ISSUED: March 23, 1999

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY Kao; Hung-Teh Hackensack NJ

Hartig; Paul R. Mahwah NJ
Branchek; Theresa Teaneck NJ

US-CL-CURRENT: 435/7.21; 435/325, 435/69.1, 530/350, 536/23.5

ABSTRACT:

The present invention provides an isolated nucleic acid molecule encoding a 5-HT.sub.2 receptor, and an isolated protein which is a human 5-HT.sub.2 receptor. The invention also provides vectors comprising DNA molecules encoding a human 5-HT.sub.2

receptor, and vectors adapted for expression of the 5-HT.sub.2 receptor in bacterial, yeast, or mammalian cells. In addition, the invention provides a DNA probe useful for detecting nucleic acid encoding the 5-HT.sub.2 receptor, a method for determining whether a ligand which is not known to be capable of binding to the 5-HT.sub.2 receptor can bind to the 5-HT.sub.2 receptor, a method for detecting the presence of 5-HT.sub.2 receptor on the surface of a cell, and a method of screening drugs to identify drugs which specifically interact with, and bind to, the 5-HT.sub.2 receptor. The invention herein also concerns an antibody directed to the human 5-HT.sub.2 receptor, such as a monoclonal antibody directed to an epitope of the 5-HT.sub.2 receptor present on the surface of a cell and having an amino acid sequence included within the amino acid sequence shown in FIGS. 2A-2G.

11 Claims, 11 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 11



73. Document ID: US 5882855 A

L13: Entry 73 of 78

File: USPT

Mar 16, 1999

US-PAT-NO: 5882855

DOCUMENT-IDENTIFIER: US 5882855 A

TITLE: DNA encoding a human dopamine D.sub.1 receptor and uses thereof

DATE-ISSUED: March 16, 1999

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Weinshank; Richard L. New York NY Hartig; Paul R. Kinnelon NJ

US-CL-CURRENT: $\underline{435}/\underline{6}$; $\underline{435}/\underline{252.3}$, $\underline{435}/\underline{320.1}$, $\underline{435}/\underline{69.1}$, $\underline{435}/\underline{7.1}$, $\underline{435}/\underline{7.2}$, $\underline{530}/\underline{350}$

ABSTRACT:

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This invention provides isolated nucleic acid molecules encoding a human dopamine D.sub.1 receptor, isolated proteins which are human dopamine D.sub.1 receptor, vectors comprising isolated nucleic acid molecules encoding a human dopamine D.sub.1 receptor, mammalian cells comprising such vectors, antibodies directed to a human dopamine D.sub.1 receptor, nucleic acid probes useful for detecting nucleic acid encoding human dopamine D.sub.1 receptor, antisense oligonucleotides complementary to any sequences of a nucleic acid molecule which encodes a human dopamine D.sub.1 receptor, pharmaceutical compounds related to human dopamine D.sub.1 receptor, and nonhuman transgenic animals which express DNA a normal or a mutant human dopamine D.sub.1 receptor. This invention further provides methods for determining ligand binding, detecting expression, drug screening, and treatment involving a human dopamine D.sub.1 receptor.

19 Claims, 10 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 10 74. Document ID: US 5763575 A

L13: Entry 74 of 78

File: USPT

Jun 9, 1998

US-PAT-NO: 5763575

DOCUMENT-IDENTIFIER: US 5763575 A

** See image for Certificate of Correction **

TITLE: Agonist and antagonist peptides of the C140 receptor

DATE-ISSUED: June 9, 1998

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Sundelin; Johan Furulund SE

Scarborough; Robert M. Belmont CA

US-CL-CURRENT: 530/327; 530/300, 530/328, 530/329, 530/330

ABSTRACT:

Nucleic acid molecules encoding the C140 cell surface receptor have been cloned and sequenced. The availability of C140 receptor DNA permits the recombinant production of the C140 receptor which can be produced on the surface of a cell, including an oocyte. The nucleic acid molecules are useful in an assay for detecting a substance which affects C140 receptor activity, either receptor agonists or antagonists. Further, the elucidation of the structure of the C140 receptor permits the design of agonist and antagonist compounds which are useful in such assays. The availability of the C140 receptor also permits production of antibodies specifically immunoreactive with one or more antigenic epitopes of the C140 receptor.

11 Claims, 20 Drawing figures Exemplary Claim Number: 1,8 Number of Drawing Sheets: 16

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Full	Title	Citation	Front	Review	Classification	Date	Reference	,q	KNNC Draw Desc

75. Document ID: US 5759804 A

L13: Entry 75 of 78 File: USPT Jun 2, 1998

US-PAT-NO: 5759804

DOCUMENT-IDENTIFIER: US 5759804 A

** See image for <u>Certificate of Correction</u> **

TITLE: Isolated nucleic acid encoding seven transmembrane receptors

DATE-ISSUED: June 2, 1998

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY Godiska; Ronald Bothell WA

Gray; Patrick W. Seattle WA Schweickart; Vicki Louise Seattle WA

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US-CL-CURRENT: 435/69.1; 435/252.3, 435/320.1, 530/350, 536/23.5

ABSTRACT:

DNA sequences encoding seven novel seven transmembrane receptors and variants thereof are disclosed as well as materials and methods for production of the same by recombinant techniques. Antibody substances specific for each of the seven transmembrane receptors are disclosed as useful for the modulation of the ligand/receptor binding reactions of the receptors.

26 Claims, 2 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 2

Full	Title Citation Fro	nt Review	Classification	Date	Reference			K	MC	Drawt Desc
	76. Document	ID: US 56	83696 A		······································	***************************************	***************************************		*********	***************************************
L13:	Entry 76 of 78	В			File	USPT		Nov	4,	1997

US-PAT-NO: 5683696

DOCUMENT-IDENTIFIER: US 5683696 A

** See image for Certificate of Correction **

TITLE: Cloning of duffy blood group antigen, gpD

DATE-ISSUED: November 4, 1997

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Pogo; Angel Oscar Pelham NY Chaudhuri; Asok Rego Park NY

US-CL-CURRENT: 424/185.1; 424/184.1, 435/69.3, 435/69.6, 435/71.1, 514/12, 514/13, 514/2, 530/300, 530/358, 530/380, 530/395

ABSTRACT:

gpD protein, the major subunit of the Duffy blood group antigenic system, has been isolated. gpD protein contains the receptor, by which P. vivax enters red cells and causes malaria. gpD has significant sequence homology with human and rabbit interleukin-8 receptors and, therefore, gpD protein likely is a new class of chemoattractant cytokines receptor. gpD protein cDNA has a quasi-total homology with a human hippocampus cDNA clone HHCMF86 and, therefore, gpD protein or a homologous protein may be present as a neuropeptide receptor in brain. gpD protein is present in all red cell progenitors and it may be a receptor for cell proliferation and/or differentiation. gpD protein cDNA identifies in human kidney a mRNA of the same size as the bone marrow. Since the kidney is not and has no potential to become an erythropoietic organ, this putative chemoattractant receptor may have essential renal functions. gpD protein has therapeutic value in the prevention of malaria and in the regulation of erythrocyte, neural and renal functions and can be combined with physiologically acceptable diluents to yield a therapeutic agent suitable for these purposes. Peptides corresponding to a portion of gpD protein that contains the receptor also have been synthesized. Such peptides have therapeutic usefulness identical to that of gpD protein. gpD protein and such peptides also have utility in the production of therapeutics, e.g., antibodies, complementary peptides, etc., which are also useful to treat malaria and regulate essential erythrocyte, neural and renal functions.

18 Claims, 9 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 7

Full Title Citation Front	Review Classification	Date Reference	,,	KMC Draw Desc

77. Document ID: US 5621079 A

L13: Entry 77 of 78

File: USPT

Apr 15, 1997

US-PAT-NO: 5621079

DOCUMENT-IDENTIFIER: US 5621079 A

TITLE: Neuropeptide Y receptor

DATE-ISSUED: April 15, 1997

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY Cascieri; Margaret A. E. Windsor N_iT Linemeyer; David L. Westfield NJ Macneil; Douglas J. Westfield NJ Shiao; Lin-Lin Avenel NJ Strader; Catherine D. Verona NJ Weinberg; David H. Westfield NJ Tan; Carina P. Metuchen NJ

US-CL-CURRENT: <u>530/350</u>; <u>435/69.1</u>

ABSTRACT:

A novel mammalian neuropeptide Y receptor and method of making the receptor are provided. The invention includes DNA encoding the receptor, the receptor, assays employing the receptor, cells expressing the receptor, antibodies which bind specifically to the receptor, RNA encoded by the DNA sequence or its complementary sequence, and single-stranded DNA with a sequence complementary to the RNA which encodes the receptor. The receptor and assays employing the receptor are useful for identifying compounds which bind to the receptor, including specific modulators of the receptor. Such compounds are useful for treating a variety of disease conditions, including obesity, diabetes, anxiety, hypertension, cocaine withdrawal, congestive heart failure, memory enhancement, cardiac and cerebral vasospasm, pheochromocytoma and ganglioneuroblastoma, and Huntington's, Alzheimer's and Parkinson's diseases.

8 Claims, 13 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 12

Full Title	Citation Fr	ont Review	Classification	Date	Paterona	,y	KMC	Drawi Desc

78. Document ID: US 5140105 A

L13: Entry 78 of 78

File: USPT

Aug 18, 1992

US-PAT-NO: 5140105

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DOCUMENT-IDENTIFIER: US 5140105 A

TITLE: Methods and materials for HIV detection

DATE-ISSUED: August 18, 1992

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Ohno; Tsuneya Ridgewood NJ

US-CL-CURRENT: <u>530/350</u>; <u>435/5</u>

ABSTRACT:

Disclosed are immunologically active polypeptides, preferably antibodies or antibody fragments, and most preferably monoclonal antibodies, which are reactive with idiotypes of antibodies to human lymphocyte T4 protein and are reactive with the HIV virion in a manner allowing for in vitro and in vivo neutralization of HIV infectivity and detection of HIV particles in biological fluids. Presently preferred embodiments comprise monoclonal anti-monoclonal-anti-human lymphocyte T4 antibodies produced by new murine hybridoma cell lines JT4C8, JT4C12, JT4C16, JT1-1F3, JT1-1F3-E5, JT1-1D7 and JT2-N15. Also disclosed are active and passive vaccination procedures.

3 Claims, 10 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 5

Full Title Citation	Front Review	Classification	Date	Reference				KAMC	Draw Desc
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Role of orexin and prostaglandin E(2) in activating histaminergic

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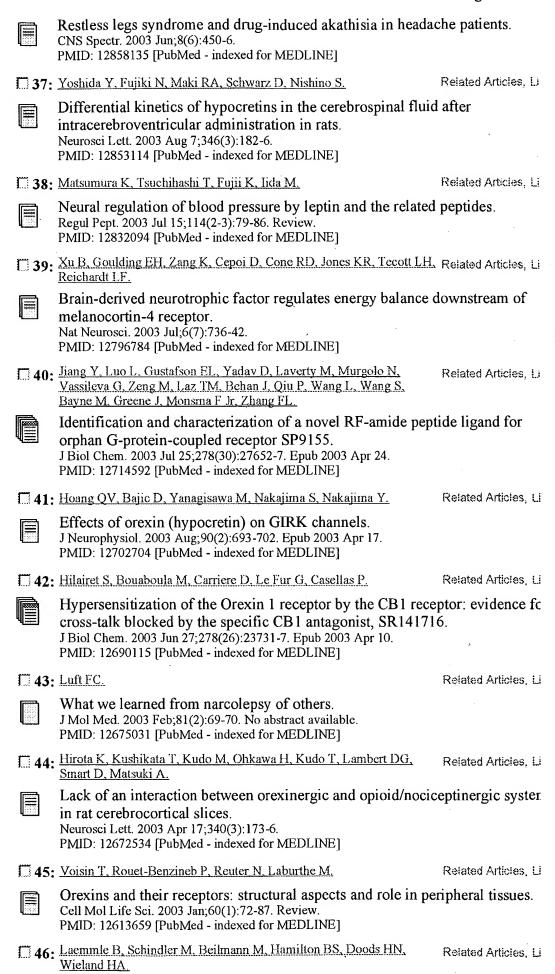
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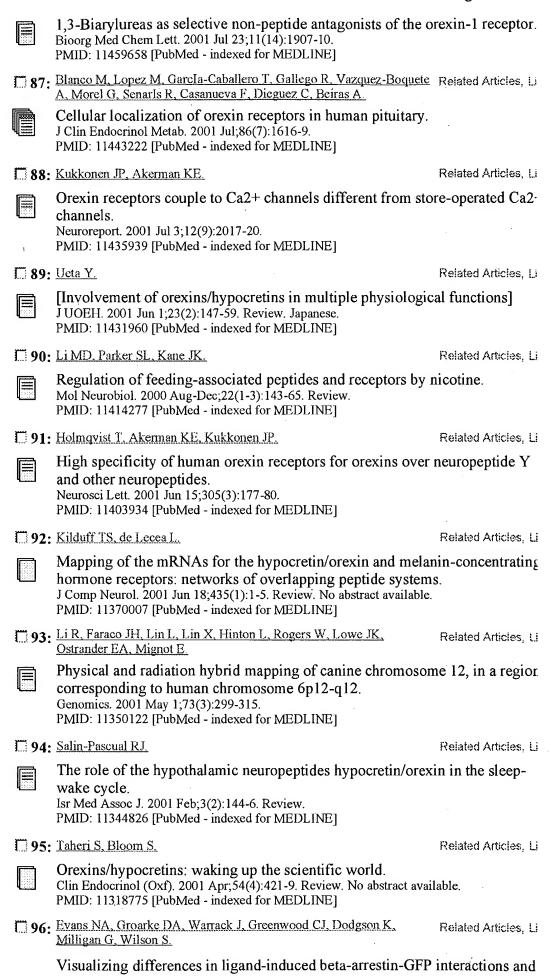
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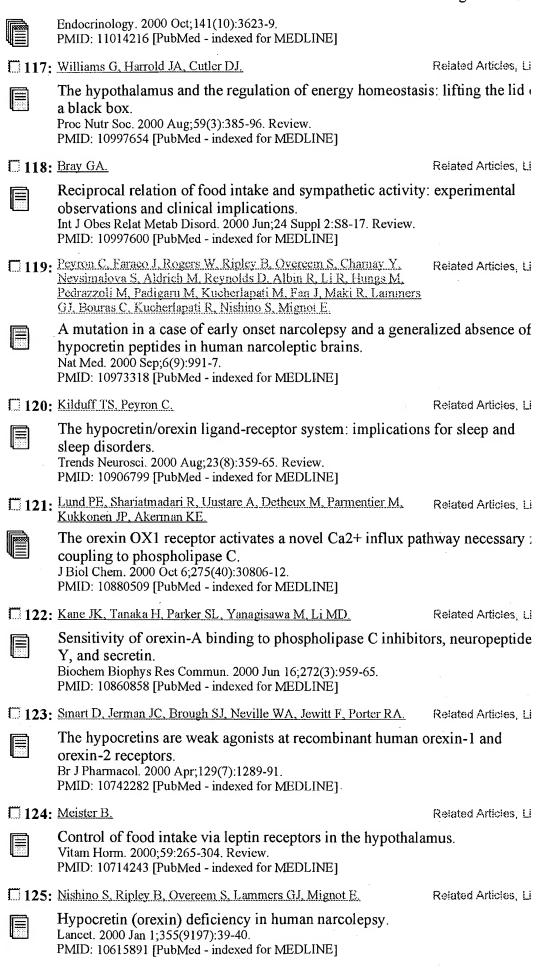
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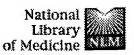
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To eat or to sleep? Orexin in the regulation of feeding and wakefulness.

Willie JT, Chemelli RM, Sinton CM, Yanagisawa M.

Howard Hughes Medical Institute, University of Texas Southwestern Medical Center at Dallas, 75390-9050, USA. willie.jon@tumora.swmed.edu

Orexin-A and orexin-B are neuropeptides originally identified as endogenous ligands for two orphan G-protein-coupled receptors. Orexin neuropeptides (als known as hypocretins) are produced by a small group of neurons in the lateral hypothalamic and perifornical areas, a region classically implicated in the cont of mammalian feeding behavior. Orexin neurons project throughout the central nervous system (CNS) to nuclei known to be important in the control of feedin sleep-wakefulness, neuroendocrine homeostasis, and autonomic regulation. orexin mRNA expression is upregulated by fasting and insulin-induced hypoglycemia. C-fos expression in orexin neurons, an indicator of neuronal activation, is positively correlated with wakefulness and negatively correlated with rapid eye movement (REM) and non-REM sleep states. Intracerebroventricular administration of orexins has been shown to significan increase food consumption, wakefulness, and locomotor activity in rodent models. Conversely, an orexin receptor antagonist inhibits food consumption. Targeted disruption of the orexin gene in mice produces a syndrome remarkab. similar to human and canine narcolepsy, a sleep disorder characterized by excessive daytime sleepiness, cataplexy, and other pathological manifestations the intrusion of REM sleep-related features into wakefulness. Furthermore, ore knockout mice are hypophagic compared with weight and age-matched littermates, suggesting a role in modulating energy metabolism. These findings suggest that the orexin neuropeptide system plays a significant role in feeding sleep-wakefulness regulation, possibly by coordinating the complex behaviora and physiologic responses of these complementary homeostatic functions.

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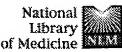
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Orexins and orexin receptors: a family of hypothalamic neuropeptides and G protein-coupled receptors that regulate feeding behavior.

Sakurai T, Amemiya A, Ishii M, Matsuzaki I, Chemelli RM, Tanaka H, Williams SC, Richardson JA, Kozlowski GP, Wilson S, Arch JR, Buckingham RE, Haynes AC, Carr SA, Annan RS, McNulty DE, Liu WS, Terrett JA, Elshourbagy NA, Bergsma DJ, Yanagisawa M.

Howard Hughes Medical Institute, Department of Molecular Genetics, Univers of Texas Southwestern Medical Center at Dallas, 75235-9050, USA.

The hypothalamus plays a central role in the integrated control of feeding and energy homeostasis. We have identified two novel neuropeptides, both derived from the same precursor by proteolytic processing, that bind and activate two closely related (previously) orphan G protein-coupled receptors. These peptide termed orexin-A and -B, have no significant structural similarities to known families of regulatory peptides. prepro-orexin mRNA and immunoreactive orexin-A are localized in neurons within and around the lateral and posterior hypothalamus in the adult rat brain. When administered centrally to rats, these peptides stimulate food consumption. prepro-orexin mRNA level is up-regulat upon fasting, suggesting a physiological role for the peptides as mediators in the central feedback mechanism that regulates feeding behavior.

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Hypocretin/orexin, sleep and narcolepsy.

Hungs M, Mignot E.

Stanford Center for Narcolepsy, Department of Psychiatry Behavioral Science Stanford University Medical Center, Palo Alto, Ca 94305-5485, USA.

The discovery that hypocretins are involved in narcolepsy, a disorder associate with excessive daytime sleepiness, cataplexy and unusually rapid transitions to rapid-eye-movement sleep, opens a new field of investigation in the area of sle control physiology. Hypocretin-1 and -2 (also called orexin-A and -B) are new discovered neuropeptides processed from a common precursor, preprohypocre Hypocretin-containing cells are located exclusively in the lateral hypothalamus with widespread projections to the entire neuroaxis. Two known receptors, Hc and Hcrtr2, have been reported. The functional significance of the hypocretin system is rapidly emerging in both animals and humans. Hypocretin abnormalities cause narcolepsy in dogs, human and mice. The role of the hypocretin system in normal sleep regulation is more uncertain. We believe hypocretin cells drive cholinergic and monoaminergic activity across the sleep cycle. Input from the suprachiasmatic nucleus to hypocretin-containing neuron may explain the occurrence of clock-dependent alertness. Other functions are suggested by pharmacological and neurochemical experiments. These include regulation of food intake, neuroendocrine function, autonomic nervous system activity and energy balance. Copyright 2001 John Wiley & Sons, Inc.

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A mutation in a case of early onset narcolepsy and a generalized absence of hypocretin peptides in human narcoleptic brains.

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Peyron C, Faraco J, Rogers W, Ripley B, Overeem S, Charnay Y, Nevsimalova S, Aldrich M, Reynolds D, Albin R, Li R, Hungs M, Pedrazz M, Padigaru M, Kucherlapati M, Fan J, Maki R, Lammers GJ, Boura's C. Kucherlapati R, Nishino S, Mignot E.

Center for Narcolepsy, Stanford University Medical School 1201 Welch Road Stanford, California 94305-5485, USA.

We explored the role of hypocretins in human narcolepsy through histopatholc of six narcolepsy brains and mutation screening of Hcrt, Hcrtr1 and Hcrtr2 in 7 patients of various human leukocyte antigen and family history status. One Hc mutation, impairing peptide trafficking and processing, was found in a single c with early onset narcolepsy. In situ hybridization of the perifornical area and peptide radioimmunoassays indicated global loss of hypocretins, without glios or signs of inflammation in all human cases examined. Although hypocretin lo do not contribute significantly to genetic predisposition, most cases of human narcolepsy are associated with a deficient hypocretin system.

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Polymorphisms in hypocretin/orexin pathway genes and narcolep

Olafsdottir BR, Rye DB, Scammell TE, Matheson JK, Stefansson K, Gulc. JR.

deCODE genetics Inc, Reykjavik, Iceland.

1: Neurology. 2001 Nov 27;57(10):1896-9.

The neuroexcitatory peptide hypocretin and its receptors are central to the pathophysiology of both human and animal models of the disease. In this study American and Icelandic patients with narcolepsy, the authors found no signific association between narcolepsy and single-nucleotide polymorphisms in the genes for hypocretin or its two known receptors, hypocretin receptor-1 and hypocretin receptor-2.

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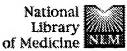
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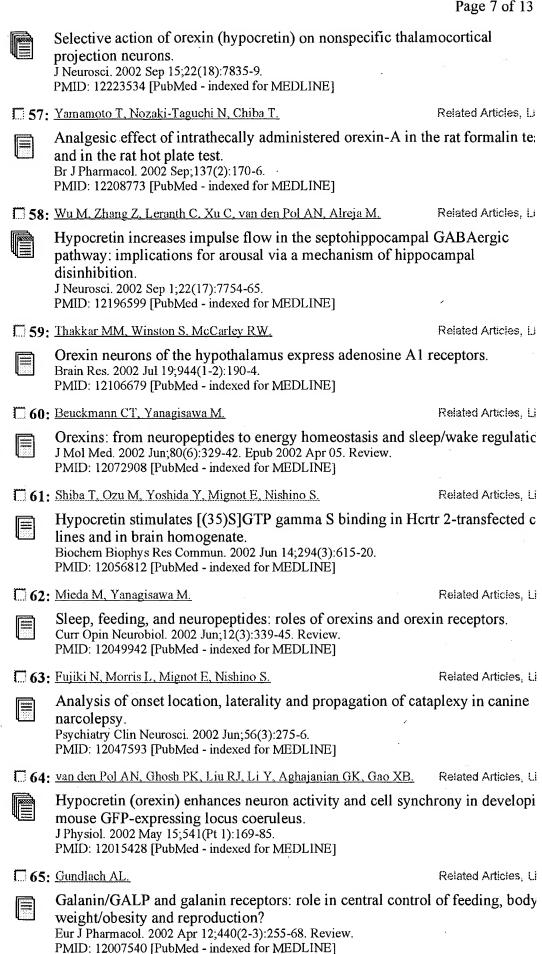
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      Institute of Basic Medical Sciences, University of Tsukuba, Tsukuba,
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      Department of Pharmacology, University of Tennessee College of Medicine, 874 Union Avenue, Memphis, TN, 38163, USA
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      Molecular Neurobiology, (August-October-December, 2000) Vol. 22. No. 1-3.
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      j.p.h.wilding@liv.ac.uk
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     University of Tsukuba, Tsukuba, Ibaraki, 305-8575, Japan Brain Research ( ***2000*** ), 873(1), 181-187
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     CODEN: BRREAP; ISSN: 0006-8993
PB
     Elsevier Science B.V.
DT
     Journal
LA
     English
RE.CNT
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               THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD
               ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 56 OF 154 CAPLUS COPYRIGHT 2004 ACS ON STN
L5
     2000:284343 CAPLUS
ΑN
DN
     133:38334
     Control of food intake via leptin ***receptors***
ΤI
                                                                  in the hypothalamus
ΑU
     Meister, Bjorn
CS
     Department of Neuroscience, Karolinska Institutet, Stockholm, S-171 77,
     Vitamins and Hormones (San Diego) ( ***2000*** ), 59, 265-304
SO
     CODEN: VIHOAQ; ISSN: 0083-6729
PR
     Academic Press
DT
     Journal; General Review
     English
LA
RE.CNT
        151
               THERE ARE 151 CITED REFERENCES AVAILABLE FOR THIS RECORD
               ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 57 OF 154 CAPLUS COPYRIGHT 2004 ACS on STN
L5
     2000:264439 CAPLUS
AN
DN
     133:26938
TI
     Structure-activity relationship studies on the novel neuropeptide
        ***orexin***
ΑU
     Asahi, Shuichi; Egashira, Shin-Ichiro; Matsuda, Masao; Iwaasa, Hisashi;
     Kanatani, Akio; Ohkubo, Mitsuru; Ihara, Masaki; Sakurai, Takeshi;
     Morishima, Hajime
CS
     Banyu Tsukuba Research Institute in collaboration with Merck Research
     Laboratories, Tsukuba, 300-2611, Japan
Peptide Science ( ***1999*** ), 36th, 37-40
SO
     CODEN: PSCIFQ; ISSN: 1344-7661
PB
     Japanese Peptide Society
     Journal
DT
     English
LA
RE.CNT
        10
               THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD
               ALL CITATIONS AVAILABLE IN THE RE FORMAT
L5
     ANSWER 58 OF 154 CAPLUS COPYRIGHT 2004 ACS on STN
AN
     2000:247790 CAPLUS
     132:277281
DN
TI
     Canine narcolepsy
ΑU
     Kadotani, Hiroshi
CS
     Sleep Disorders Cent., Stanford Univ., 1201 Welch Road, MSLS, P126,
     Stanford, CA, 94305, USA
No no Kagaku ( ***2000***
                                    ), 22(4), 465-468
SO
     CODEN: NNOKFZ; ISSN: 1343-4144
PB
     Seiwa Shoten
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Journal; General Review
DT
LA
     Japanese
L5
     ANSWER 59 OF 154 CAPLUS COPYRIGHT 2004 ACS on STN
     1999:795994 CAPLUS
ΑN
     132:31744
DN
     Gene probes used for genetic profiling in healthcare screening and
TT
     planning
IN
     Roberts, Gareth Wyn
     Genostic Pharma Ltd., UK
PA
     PCT Int. Appl., 745 pp.
so
     CODEN: PIXXD2
DT
     Patent
     English
LA
FAN.CNT 2
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                                     DATE
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     wo 9964627
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                                     19980606
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      GB 1998-13291
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      132:31743
DN
      Gene probes used for genetic profiling in healthcare screening and
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      Roberts, Gareth Wyn
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      Genostic Pharma Limited, UK
SO
      PCT Int. Appl., 149 pp.
      CODEN: PIXXD2
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FAN.CNT 2
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               TM,
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                        TJ,
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                                                   CA 1999-2330929
     CA 2330929
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     AU 9941586
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                                                  AU 1999-41586
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     AU 766544
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                              A1
                                     19991230
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                              A1 ·
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JP 2003528564
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                                                                           19990604
     US 2003198970
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                             Α1
                                                                           20020729
PRAI GB 1998-12098
                                    19980606
                             Α
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L5
     ANSWER 61 OF 154 CAPLUS COPYRIGHT 2004 ACS on STN
ΑN
     1999:571534 CAPLUS
     131:284870
DN
     Narcolepsy in ***orexin***
                                        knockout mice: molecular genetics of sleep
TI
     regulation
ΑU
     Chemelli, Richard M.; Willie, Jon T.; Sinton, Christopher M.; Elmquist,
     Joel K.; Scammell, Thomas; Lee, Charlotte; Richardson, James A.; Williams,
     Clay; Xiong, Yumei; Kisanuki, Yaz; Fitch, Thomas E.; Nakazato,
     Masamitsu; Hammer, Robert E.; Saper, Clifford B.; Yanagisawa, Masashi
     Howard Hughes Medical Institute Department of Molecular Genetics
CS
     Department of Pediatrics, University of Texas Southwestern Medical Center
     at Dallas, Dallas, TX, 75235-9050, USA
Cell (Cambridge, Massachusetts) ( ***1999*** ), 98(4), 437-451
SO
     CODEN: CELLB5; ISSN: 0092-8674
     Cell Press
PB
DT
     Journal
     English
LA
RE.CNT
        54
               THERE ARE 54 CITED REFERENCES AVAILABLE FOR THIS RECORD
               ALL CITATIONS AVAILABLE IN THE RE FORMAT
L5
     ANSWER 62 OF 154 CAPLUS COPYRIGHT 2004 ACS on STN
     1999:566074 CAPLUS
ΑN
     131:194807
DN
     Insulinotropic N-terminally truncated GLP-1 lipophilic derivatives with
TI
     protracted action
     Knudsen, Liselotte Bjerre; Huusfeldt, Per Olaf
IN
     Novo Nordisk A/s, Den.
PΑ
SO
     PCT Int. Appl., 50 pp.
     CODEN: PIXXD2
DT
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IΑ
FAN.CNT 12
     PATENT NO.
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PΙ
     wo 9943705
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              AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP,
              KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN,
              MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM,
              TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ,
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A1 20001206 EP 1999-906075 19990225
     AU 9926105
                                                                           19990225 <--
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                                                                           19990225 <--
              AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE, FI
508162 T2 20020319 JP 2000-533455 19990225
     JP 2002508162 T2
PRAI DK 1998-264
                                    19980227
     DK 1998-509
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     WO 1999-DK81
                                   19990225
os
     MARPAT 131:194807
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               THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD
               ALL CITATIONS AVAILABLE IN THE RE FORMAT
L5
     ANSWER 63 OF 154 CAPLUS COPYRIGHT 2004 ACS on STN
     1999:404816 CAPLUS
ΑN
     131:68557
DN
     Methods of treatment of behavioral and metabolic disorders using novel
TI
     ligands of the neuropeptide
                                       ***receptor***
                                                        HGFAN72 and agonists or
     antagonists thereof
ΙN
     Hagan, James J.; Kennett, Guy A.; Patel, Saraswati R.; Piper, David;
     Smith, Martin I.; Terrett, Jonathan A.; Upton, Neil
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SmithKline Beecham PLC, UK
50
     PCT Int. Appl., 68 pp.
     CODEN: PIXXD2
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     PATENT NO.
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     wo 9930670
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                                  20040303
             BE, CH, DE, DK, FR, GB, IT, LI, NL
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     us 2003087801
                                              us 1998-211823
                           В2
     us 6664229
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                           T2
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     US 1997-69785P
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                                  19971216
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                           W
                                 19981215
     ANSWER 64 OF 154 CAPLUS COPYRIGHT 2004 ACS ON STN
     1999:90461
ΑN
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DN
     130:149584
ΤI
     sequence and therapeutic applications for cDNA clone my1 that encodes a
                                             ***receptor***
             ***human***
                           7-transmembrane
ΙN
     Yanagisawa, Masashi
PΑ
     SmithKline Beecham Corporation, USA
S0
     Eur. Pat. Appl., 23 pp.
     CODEN: EPXXDW
DT
     Patent
    English
LA
FAN.CNT f 1
     PATENT NO.
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                                                                       DATE
PΙ
    EP 893498
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     US 6166193
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     CA 2238655
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        11178588
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                                              JP 1998-242457
                                                                       19980724 <--
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        2003083466
                           Α1
                                 20030501
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                                                                       20021028
PRAI US
       1997-53790P
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                                  19970725
    US 1998-119788
                           Α3
                                 19980721
    us 2000-676625
                           Α1
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L5
    ANSWER 65 OF 154 CAPLUS COPYRIGHT 2004 ACS ON STN
    1998:414800 CAPLUS
AΝ
     129:77033
DN
TI
    Novel ligands for the G protein-coupled neuropeptide
                                                              ***receptor***
    HFGAN72 and cDNAs encoding them
ΙN
    Bergsma, Derk J.; Brooks, David P.; Gellai, Miklos; Yanagisawa, Masashi;
    Wilson, Shelagh
PΑ
    Smithkline Beecham Corp., USA; Smithkline Beecham Plc
SO
    Eur. Pat. Appl., 35 pp.
    CODEN: EPXXDW
    Patent
    English
LA
FAN.CNT<sup>-</sup>2
    PATENT NO.
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                                 DATE
                                              APPLICATION NO.
                                                                       DATE
PΙ
    EP 849361
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    US 6001963
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                                              US 1997-938548
                                                                       19970926 <--
    US 6309854
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PRAI US 1996-33604P
                           Р
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    ANSWER 66 OF 154 CAPLUS COPYRIGHT 2004 ACS on STN
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PA

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ΑN
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     129:63410
                                ***Orexin***
                                                    ***Receptors*** : A Family of
TI
        ***Orexins***
                         and
     Hypothalamic Neuropeptides and G Protein-Coupled ***Receptors*** t
Regulate Feeding Behavior. [Erratum to document cited in CA128:290571]
Sakurai, Takeshi; Amemiya, Akira; Ishii, Makoto; Matsuzaki, Ichiyo;
Chemelli, Richard M.; Tanaka, Hirokazu; Williams, S. Clay; Richardson,
James A.; Kozlowski, Gerald P.; Wilson, Shelagh; Arch, Jonathan R. S.;
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     Buckingham, Robin E.; Haynes, Andrea C.; Carr, Steven A.; Annan, Roland
     S.; McNulty, Dean E.; Liu, Wu-Schyong; Terrett, Jonathan A.; Elshourbagy,
     Nabil A.; Bergsma, Derk J.; Yanagisawa, Masashi
CS
     Howard Hughes Medical Institute, Department of Molecular Genetics,
     University of Texas Southwestern Medical Center at Dallas, Dallas, TX,
     75235-9050, USA
     Cell (Cambridge, Massachusetts) ( ***1998*** ), 92(5), No pp. Given
50
     CODEN: CELLB5; ISSN: 0092-8674
PB
     Cell Press
DT
     Journal
     English
LA
L5
     ANSWER 67 OF 154 CEN COPYRIGHT 2001 ACS ON STN
ΑN
     1999:2064 CEN
TI
     Appetite-regulating peptide linked to sleep disorder
     Chemical & Engineering News, ( ***9 Aug 1999*** ) Vol. 77, No. 32, pp.
S0
     CODEN: CENEAR, ISSN: 0009-2347.
PB
     American Chemical Society
     English
LA
WC
L5
     ANSWER 68 OF 154 CIN COPYRIGHT 2004 ACS on STN
ΑN
     28(34):34208s CIN
TI
     Appetite-regulating peptide linked to sleep disorder
     Chem. Mark. Rep., 9 Aug 1999 (19990809), 256(6), p. 27. ISSN: 0009-2347;
S0
     CODEN: CMREF6.
     English
LA
L5
      ANSWER 69 OF 154 DGENE COPYRIGHT 2004 The Thomson Corp on STN
ΑN
      AAB07426 Protein
                                  DGENE
      Novel polynucleotide encoding G protein coupled
TI
                                                              ***receptor***
      useful for producing recombinant cell lines for discovering therapeutic
      agents that modulate the ***receptor***
                                                        activity -
      Zastawny R L
IN
       (ALLX)
                   ALLELIX BIOPHARMACEUTICALS INC.
PA
        ***CA 2284857
ΡI
                          A1 20000416
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      CA 1999-2284857
ΑI
                              19991015
PRAI
      US 1998-104514
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      US 1998-173565
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DT
      Patent
LA
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os
      2000-491457 [44]
CR
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DESC
      Amino acid sequence of a
                                   ***human***
                                                     Α4
                                                           ***receptor***
      polypeptide.
L5
      ANSWER 70 OF 154 DGENE COPYRIGHT 2004 The Thomson Corp on STN
                                 DGENE

***receptor***
ΑN
      AAB21212 peptide
                   ***orexin***
TI
                                                          antagonist for treating
      disorders associated with neuronal degeneration resulting from ischemic
      events, nausea and vomiting, irritable bowel syndrome or other conditions
      associated with visceral pain -
ΙN
      Irving E A; Sanger G J
PA
                    SMITHKLINE BEECHAM PLC.
        ***WO 2000047284 A2 20000817
                                                           10p***
PΙ
      WO 2000-EP1147
ΑI
                              20000210
PRAI
      GB 1999-3265
                              19990212
      GB 1999-3278
                              19990212
      GB 1999-3282
                              19990212
      GB 1999-3284
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      GB 1999-6061
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DT
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LA
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วร
      2000-532977 [48]
        ***Human***
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DESC
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L5
      ANSWER 71 OF 154 DGENE COPYRIGHT 2004 The Thomson Corp on STN
ΑN
      AAB21211 peptide
                                DGENE
      New N-(1,5-naphthyridin-4-yl)-N'-phenylurea derivatives, used to treat
ΤI
      e.g. obesity, diabetes, sleep disorders, pain, migraine, heart and lung
      diseases, depression, schizophrenia, addictions and sexual dysfunction,
            ***orexin***
                           -1 antagonists ·
      Coulton S; Johns A; Porter Ř A
ΙN
PΑ
      (SMIK)
                   SMITHKLINE BEECHAM PLC.
        ***WO 2000047580 A2 20000817
PΙ
                                                        28p***
AΙ
      WO 2000-EP1142
                             20000210
      GB 1999-3241
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                             19990212
      GB 1999-26441
                             19991108
DΤ
      Patent
      English
ŀΑ
      2000-515054 [46]
กร
        ***Human***
                         ***orexin*** -A.
DESC
L5
      ANSWER 72 OF 154 DGENE COPYRIGHT 2004 The Thomson Corp on STN
      AAB21210 peptide
ΑN
                                DGENE
ΤI
      Use of new and known N-phenyl-N'-(4-quinolinyl)urea derivatives, used to
      treat e.g. obesity, diabetes, sleep disorders, pain, migraine, heart and lung disorders, depression and addictions are ***orexin*** -1
      lung disorders, depression and addictions are
      antagonists ·
ΙN
      Coulton S; Johns A; Porter R A
        MIK) SMITHKLINE BEECHAM PLC. ***WO 2000047577 A1 20000817
PA
      (SMIK)
PΙ
                                                        45p***
      WO 2000-EP1150
AΙ
                             20000210
PRAI
      GB 1999-3266
                             19990212
      GB 1999-26430
                             19991108
DT
      Patent
LΑ
      English
วร
      2000-515053 [46]
DESC
        ***Human***
                         ***orexin*** -A.
      ANSWER 73 OF 154 DGENE COPYRIGHT 2004 The Thomson Corp on STN
L5
      AAB21209 peptide
                                DGENE
      New N-(quinolin-4-yl)-acrylamide derivatives, used to treat e.g. obesity,
ΤI
      diabetes, prolactinoma, dwarfism, sleep disorders, narcolepsy, insomnia, heart and lung diseases and depression, are ***orexin*** -1
      antagonists -
ΙN
      Johns A; Porter R A
                   SMITHKLINE BEECHAM PLC.
PA
      (SMIK)
        ***WO 2000047576 A1 20000817
PΙ
                                                        29p***
     WO 2000-EP1148
٩I
                             20000210
PRAI
     GB 1999-3287
                             19990212
      GB 1999-3288
                             19990212
ΣT
      Patent
      English
∟A
วร
      2000-506092 [45]
                         ***orexin*** -A.
DESC
        ***Human***
_5
     ANSWER 74 OF 154
                         DGENE COPYRIGHT 2004 The Thomson Corp on STN
NΑ
     AAA57846 DNA
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ΓI
     Novel polynucleotide encoding G protein coupled
                                                           ***receptor***
      useful for producing recombinant cell lines for discovering therapeutic
                                   ***receptor***
      agents that modulate the
                                                      activity -
ľΝ
     Zastawny R L
РΑ
      (ALLX)
                   ALLELIX BIOPHARMACEUTICALS INC.
PΙ
        ***CA 2284857
                          A1 20000416
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Ί
      CA 1999-2284857
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     US 1998-104514
                             19981016
     US 1998-173565
                             19981016
TC
     Patent
_A
     English
วร
     2000-491457 [44]
DESC
     PCR primer for DNA encoding a
                                       ***human***
                                                        Α4
                                                              ***receptor***
     polypeptide.
     ANSWER 75 OF 154 DGENE COPYRIGHT 2004 The Thomson Corp on STN
١N
     AAA57845 DNA
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     Novel polynucleotide encoding G protein coupled
ΓI
                                                            ***receptor***
     useful for producing recombinant cell lines for discovering therapeutic
     agents that modulate the
                                   ***receptor***
                                                      activity -
N
     Zastawny R L
                  ALLELIX BIOPHARMACEUTICALS INC.
Α
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     PCR primer for DNA encoding a
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     ANSWER 76 OF 154 DGENE COPYRIGHT 2004 The Thomson Corp on STN
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     Novel polynucleotide encoding G protein coupled
     useful for producing recombinant cell lines for discovering therapeutic
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      Pharmacological treatment of obesity: therapeutic strategies.
TI
      Kordik C P; Reitz A B
ΑU
CS
      Johnson+Johnson
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      Spring House, Pa., USA
      J.Med.Chem. (42, No. 2, 181-201, 1999) 3 Fig. 3 Tab. 239 Ref.
S<sub>0</sub>
                          ISSN: 0022-2623
      CODEN: JMCMAR
      Drug Discovery Division, The R.W. Johnson Pharmaceutical Research
ΑV
      Institute, Spring House, Pennsylvania 19477, U.S.A. (e-mail:
      ckordik@prius.jnj.com).
      English
LA
DT
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TT
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     Redline S.; Tishler P.V.
Dr. S. Redline, Rainbow Babies and Childrens' Hosp., Case Western Reserve
ΑU
     University, 11100 Euclid Avenue, Cleveland, OH 44106-6003, United States.
     sxr15@po.cwru.edu
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     Refs: 116
      ISSN: 1087-0792 CODEN: SMREFC
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     United Kingdom
     Journal; General Review
DT
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              Neurology and Neurosurgery
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      ANSWER 85 OF 154 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS
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ΑN
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ΤI
      GENETICKE ASPEKTY PORUCH SPANKU.
      Sonka K.; Nevsimalova S.
ΑU
      Dr. K. Sonka, Neurologicka Klinika, I Lekarska Fakulta, UK a VFN,
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      Katerinska 30, 120 00 Praha 2, Czech Republic. ksonka@lf1.cuni.cz
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      ISSN: 1211-7579 CODEN: PCHIF7
      Czech Republic
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ΑN
      'Neuropeptides 2000'10(th) Annual Meeting of the European Neuropeptide Club, Innsbruck, Austria, May 10-13, 2000, Neuropeptide Antagonists, From Molecular Biology to ***Receptors*** and Clinical Applications,
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      Refs: 0
      ISSN: 0167-0115 CODEN: REPPDY
      s 0167-0115(00)00191-9
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      RESERVED. on STN 2000351213 EMBASE
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      Obesity: Molecular bases of a multifactorial problem.
     Palou A.; Serra F.; Bonet M.L.; Pico C.
A. Palou, Dept. Biol. Fonam. Cien. de la Salut, Universitat de les Illes
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      Palma de Mallorca, Spain. dbfapo0@ps.uib.es
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SO
      Refs: 196
      ISSN: 1436-6207 CODEN: EJNUFZ
      Germany
      Journal; General Review
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ISSN: 1354-3776 CODEN: EOTPEG
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           Halford J.C.G.; Blundell J.E.
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           Dr. J.C.G. Halford, Department of Psychology, University of Liverpool,
 CS
           Liverpool L69 3BX, United Kingdom
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           Refs: 151
           ISSN: 0071-786X CODEN: FAZMAE
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ΑU
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          Dr. J.C.G. Halford, Department of Psychology, Eleanor Rathbone Building,
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          University of Liverpool, Liverpool L69 7ZA, United Kingdom. j.c.g.halford@liverpool.ac.uk
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          2000136660 EMBASE
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TI
          Schwartz M.W.; Woods S.C.; Porte D. Jr.; Seeley R.J.; Baskin D.G.
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          ISSN: 0028-0836 CODEN: NATUAS
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          United Kingdom
          Journal; General Review
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      Smith G.P.
      G.P. Smith, Department of Psychiatry, Joan/Sanford I. Weill Med. College,
CS
      New York-Presbyterian Hospital, 21 Bloomingdale Road, White Plains, NY
      10605, United States. gpsmith@med.cornell.edu
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ΑN
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FS
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TI
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     s 0006-8993(99)01961-7
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CS
     Dr. M.L. Schubert, McGuire VAMC, Code 111N, Gastroenterology Division,
     1201 Broad Rock Boulevard, Richmond, VA 23249, United States.
     Mitchell.Schubert@med.va.gov
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SO
     Refs: 44
     ISSN: 0267-1379 CODEN: COGAEK
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AN
        ***Orexins***
                                                  ***receptors*** : Implication in
TI
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                                ***orexin***
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      Sakurai T.
ΑU
CS
      Dr. T. Sakurai, Institute of Basic Medical Sciences, University of
      Tsukuba, Tsukuba, Ibaraki 305-8575, Japan. stakeshi@md.tsukuba.ac.jp
S<sub>0</sub>
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      Refs: 26
      ISSN: 0167-0115 CODEN: REPPDY
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      ANSWER 97 OF 154 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS
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AN
TI
      Narcolepsy and the
                            ***hypocretin***
                                                   ***receptor***
                                                                     2 gene.
     Aldrich M.S.; Reynoldst P.R.
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     Medicine, P.O. Box 100244, Gainesville, FL 32610-0244, United States.
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CY
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       ***Orexins***
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006
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L5
       ANSWER 100 OF 154 Elsevier BIOBASE COPYRIGHT 2004 Elsevier Science B.V.
       on STN
AN
       2000164552
                       ESBIOBASE
       Reciprocal relation of food intake and sympathetic activity: Experimental
TI
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ΑU
       Bray G.A.
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SO
       101 reference(s)
       CODEN: IJOBDP ISSN: 0307-0565
DT
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       United Kingdom
CY
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SO
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       (24 ref.)
       Published by: CABI Publishing, CAB International. Address: Wallingford, Oxon OX10 8DE, UK. Telephone: +44 (1491) 832111. Fax: +44 (1491)
       829198. Email: publishing@cabi.org Web: http://nutrition.cabweb.org and
       www.nutsoc.org.uk
       ISSN: 0007-1145
DT
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GenBank VERSION (VER):
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SEQUENCE LENGTH (SQL):
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MOLECULE TYPE (CI):
DIVISION CODE (CI):
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DATE (DATE).
                            6 Feb 2001
DEFINITION (DEF):
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(HCRTR2) genes, complete cds.
                                              ***hypocretin***
                                                                       ***receptor*** -2
SOURCE:
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   AUTHOR (AU):
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                            Charnay,Y.; Nevsimalova,S.; Aldrich,M.; Reynolds,D.; Albin,R.; Li,R.; Hungs,M.; Pedrazzoli,M.; Padigaru,M.; Kucherlapati,M.; Fan,J.; Maki,R.; Lammers,G.J.; Bouras,C.; Kucherlapati,R.; Nishino,S.; Mignot,E. A mutation in a case of early onset narcolepsy and a generalized absence of ***hypocretin*** peptides in ***hyman***
   TITLE (TI):
                                                                                peptides in
                              ***human***
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   JOURNAL (SO):
                            Nat. Med., 6 (9), 991-997 ( ***2000*** )
   OTHER SOURCE (OS):
                            CA 133:348631
REFERENCE:
                               (bases 1 to 4610)
   AUTHOR (AU):
                            Faraco, J.; Rogers, W.; Overeem, S.; Li, R.; Mignot, E.
   TITLE (TI):
                            Direct Submission
   JOURNAL (SO):
                            Submitted (05-NOV-1999) Center for Narcolepsy Research,
                            Department of Psychiatry, Stanford University Medical
                            Center, 1201 Welch Road, MSLS Bldg. P112, Stanford, CA
                            94305-5485, USA
FEATURES (FEAT):
 Feature Key
                       Location
                                                   Qualifier
                   3452..3667
source
                                               /organism="Homo sapiens"
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/db-xref="taxon:9606"

		/note="amplify at 58 degrees,
		R2-ex4-F: 5`-AAGGTAAATATGCACTTTGAAGAA-3',
		R2-ex4-R: 5`-AAGCACAGACATAATATTTGGAAG-3'"
source	10761291	organism="Homo sapiens"
		/db-xref="taxon:9606"
		<pre>/note="amplify at 58 degrees, R1-ex4-F:</pre>
		5`-CTGTCTGTCATGGTGGCTGTATGG-3',
		R1-ex4-R:
	2000 4210	5`-CTCTCTTTGGTTGCAGCCAAGATG-3'"
source	39894210	/organism="Homo sapiens" /db-xref="taxon:9606"
		/note="amplify at 58 degrees,
		R2-ex6-F:
	·	5`-GAGTCAGACCATCCTCTACCAATA-3',
		R2-ex6-R: 5`-ACTCACATAGCACCTAAACTCCTC-3'"
source	16191840	/organism="Homo sapiens"
		/db-xref="taxon:9606"
•		/note="amplify at 58 degrees, R1-ex6-F:
		5`-TGGGCAGTAGGAACTCTTGCACT-3',
	·	R1-ex6-R:
source	453731	5`-CAGGTACATCCTCACCCACCATC-3'"
3001 CE	4331.731	/organism="Homo sapiens" /db-xref="taxon:9606"
4		/note="amplify at 58 degrees,
		R1-ex2-F:
		5`-GAAGGGGTTGTGTGGGAAGAG-3', R1-ex2-R:
	•	5`-ACACTTCAGGGGTCATGAGCCA-3'"
source	28283107	/organism="Homo sapiens"
		/db-xref="taxon:9606"
		<pre>/note="amplify at 58 degrees, R2-ex2-F:</pre>
		5`-TGACAGTGTTTCCTCACCAATACC-3',
	,	R2-ex2-R:
source	36683988	5`-TCCTTCAGTTTGTCAATGCCTTAG-3'" /organism="Homo sapiens"
	30003300	/db-xref="taxon:9606"
		/note="amplify at 58 degrees,
		R2-ex5-F:
		5`-TCTGGAAGCCTTTCCTTACTGTG-3', R2-ex5-R:
		5`-CTTAAAGGCTGTTCGCCTTACC-3'"
source	12921618	/organism="Homo sapiens"
		/db-xref="taxon:9606" /note="amplify at 58 degrees.
		R1-ex5-F:
		5`-TTTTATCCTTTTGCCCATCTCCAC-3',
	,	R1-ex5-R: 5`-GGAGGCTCAGAGAAGAGAAATGGC-3'"
source	7321075	/organism="Homo sapiens"
		/db-xref="taxon:9606"
		/note="amplify at 58 degrees,
		R1-ex3-F: 5`-CGTCAGCCTCACTCACCTACT-3',
		R1-ex3-R:
source	31083451	5`-TGGTAGGAGCCAGTCTAGGGTGTC-3'"
Sour Ce	31063431	/organism="Homo sapiens" /db-xref="taxon:9606"
		/note="amplify at 58 degrees,
		R2-ex-3-F:
		<pre>5`-TTTTGGCAGCTTTGAATTTGCTTA-3', R2-ex3-R:</pre>
		5`-TCAAGTTGGTTTTCATGCTCTTGC-3'"
source	18412191	/organism="Homo sapiens"
		/db-xref="taxon:9606"
		/note="amplify at 58 degrees, R1-ex7-F:
		5`-CTCATAGGCAGCTTGGCTGGAG-3',
		R1-ex7-R:
source	42114610	5`-CCAGAGTCACACAGGCAGAAACC-3'" /organism="Homo sapiens"
		, or garrisme home saptens

```
/db-xref="taxon:9606"
                                                   /note="amplify at 58 degrees,
                                                   R2-ex7-F:
                                                   5 -CCCATCTTTGCAAAATATTACACC-3',
                                                   R2-ex7-R:
                                                     -CCTGAAATAAGCTCAATTGAAGGT-3"
                                                   organism="Homo sapiens"
/db-xref="taxon:9606"
source
                    1..452
                                                   /note="amplify at 58 degrees,
                                                   R1-ex1-F:
                                                   5`-CCTCCACCAATTTCATGACTGTGA-3',
                                                   R1-ex1-R:
                                                   5 -CAGAGCCACACCCATCCTAGTTCT-3'"
                                                   /organism="Homo sapiens'
/db-xref="taxon:9606"
source
                    2192..2827
                                                   /note="amplify at 58 degrees,
                                                   R2-ex1-F:
                                                   5`-CTTCAGCTTCAGCTCTCCCTCA-3',
                                                   R2-ex1-R:
                                                   5`-GAGCAGCGACCTCTTTGTTTGC-3'"
                    1..4610
                                                   /organism="Homo sapiens"
source
                                                   /db-xref="taxon:9606"
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   join(AF202078.1:1..452,gap(),AF202079.1:1..279,gap(),
AF202080.1:1..344,gap(),AF202081.1:1..216,gap(),AF202082.1:1..327,
gap(),AF202083.1:1..222,gap(),AF202084.1:1..351,gap(),
AF202085.1:1..636,gap(),AF202086.1:1..280,gap(),AF202087.1:1..344,
    gap(), AF202088.1:1..216, gap(), AF202089.1:1..321, gap(),
    AF202090.1:1..222,gap(),AF202091.1:1..400)
L5
      ANSWER 103 OF 154
                                 GENBANK RTM. COPYRIGHT 2004 on STN
LOCUS (LOC):
                              LMFL6071
                                              GenBank (R)
GenBank ACC. NO. (GBN): AL583933
GenBank VERSION (VER):
                              AL583933.1 GI:13122223
CAS REGISTRY NO. (RN):
                              324731-47-7
SEQUENCE LENGTH (SQL):
                              34156
MOLECULE TYPE (CI):
                              DNA; linear
DIVISION CODE (CI):
                              Invertebrates
DATE (DATE):
                              22 Feb 2001
DEFINITION (DEF):
                              Leishmania major Friedlin chromosome 5 cosmid L6071,
                              PREFINAL.
SOURCE:
                              Leishmania major.
 ORGANISM (ORGN):
                              Leishmania major
                              Eukaryota; Euglenozoa; Kinetoplastida;
                              Trypanosomatidae; Leishmania
NUCLEIC ACID COUNT (NA): 6951 a
                                          10087 c
                                                                    6175 t
                                                                               504 others
COMMENT:
      see http://www.ebi.ac.uk/parasites/leish.html
      Details of leishmania sequencing at the Sanger Centre are available
      on the World Wide Web.
      see http://www.sanger.ac.uk/Projects/L_major/
     CDS are numbered using the following system eg L6071.01. L6071 (cosmid name), .01 (first CDS)

To make the cosmid library Leishmania major Friedlin DNA was partially digested with Sau3AI prior to cloning into BamHI site of the cosmid shuttle vector CLHYG (Ryan et al. 1993 Gene
      131:145-150). The sequence of the packaged vector was determined by
      Peter Myler and Ken Stuart at Seattle Biomedical Research
      Institute, and is available as accession number U59231.
      The more significant matches with motifs in the PROSITE database
      are also included but some of these may be fortuitous. The length
      in codons is given for each CDS.
     Usually the highest scoring match found by fasta -o is given for CDS which show significant similarity to other CDS in the database.
      Gene prediction is done using:
      (1)
     the FramePlot program of Bibb et al.
     Gene 30:157-166(1984) as implemented
     at http://www.nih.go.jp/
     jun/cgi-bin/frameplot.pl. (2)
     codon preference based on the codon usage table for Leishmania at
     http://www.kazusa.or.jp/codon/
```

(3)

```
the Hexamer program which was written by Richard Durbin as an
       integral part of the ACEDB-based analysis tools for the C.elegans
       Genome Sequencing Project. The program calculates the
       log-likelihood score for a given DNA segment based on the frequency
      of 6-mers, normalised for the base-pair composition of the genome. The program was trained using a fasta file of confirmed Leismania major coding sequences (CDS), i.e. from ATG start codon to the stop
       codon.
                    We may not have predicted the correct initiation codon.
       Where possible we choose an initiation codon (atg) which is
       preceded by a stretch of pyrimidines or part of a Kozak sequence. If this cannot be identified we choose the most upstream initiation
       codon. Transmembrane domains were predicted as implemented at the
       TMHMM server: http://www.cbs.dtu.dk/services/TMHMM-1.0/
       IMPORTANT: This sequence MAY NOT be the entire insert of the
       sequenced clone. It may be shorter because we only sequence
      overlapping sections once, or longer, because we arrange for a small overlap between neighbouring submissions. Cosmid L6071 is overlapped at the 5' end by L4370 (not sequenced), contains the 'right end' sequences for PACS P719 (AL161399) and P108 (AL160498). P719 links into L7758 (AL352980), while P108 links into L2267 (AL357593). Cosmid L6071 is overlapped in the middle by L6812.2 (to
       be sequenced).
REFERENCE:
                                    (bases 1 to 34156)
    AUTHOR (AU):
                                 Ivens,A.C.; Lewis,S.M.; Bagherzadeh,A.; Zhang,L.;
                                 Chan, H.M.; Smith, D.F.
                                 A physical map of the Leishmania major Friedlin genome
    TITLE (TI):
                                                 8 (2), 135-145 ( ***1998*** )
    JOURNAL (SO):
                                 Genome Res.,
                                 CA 128:266833
2 (bases 1 to 34156)
    OTHER SOURCE (OS):
REFERENCE:
    AUTHOR (AU):
                                 Zimmermann, W.; Ivens, A.C.; Quail, M.; Rajandream, M.A.;
                                 Barrell, B.G.
    TITLE (TI):
                                 Direct Submission
    JOURNAL (SO):
                                 Submitted (20-FEB-2001) European Leishmania major
                                 Friedlin genome sequencing project, Sanger Centre, The
                                Wellcome Trust Genome Campus, Hinxton, Cambridge CB10
                                 1SA, (E-mail: barrell@sanger.ac.uk) and Agowa GmbH,
                                 Glienickerweg 185, D-12489, Berlin, Germany
FEATURES (FEAT):
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                           Location
                                                            Qualifier
  =======+==
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source
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                                                       /db-xref="taxon:5664"
                                                       /chromosome="5"
                                                       /clone="cosmid L6071"
                                                      /clone= cosmid LoU/I
/note="poly-pyrimidine tract"
/note="poly-pyrimidine tract"
/note="(ctc)6"
/note="poly-pyrimidine tract"
/note="poly-pyrimidine tract"
/note="(atgt)4"
/note="(acgtatat)4"
repeat-region
                      371..386
                      436..456
repeat-region
                     complement(569..604)
complement(578..595)
complement(666..697)
repeat-region
repeat-region
repeat-region
repeat-region
                      734..751
                      complement(953..968)
repeat-region
                                                       /note="(acatatat)4"
repeat-region
                      complement(969..1000)
                                                       /note="poly-pyrimidine tract"
repeat-region
                      complement(1095..1112)
                                                      /note="poly-pyrimidine tract"
/note="poly-pyrimidine tract"
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repeat-region
                      complement(1143..1157)
repeat-region
                      complement(1303..1322)
gene
                      complement(1575..3299)
                                                      /gene="L6071.01"
/note="L6071.01, len = 573 aa,
                      complement(1575..3299)
                                                       possible monocarboxylate
                                                      transporter protein; weak
                                                       (sub-threshold) Pfam match to
                                                      entry PF01587, Monocarboxylate
                                                      transporter; MCTs catalyse the
                                                      proton linked transport of lactic
                                                      acid, pyruvate and ketone bodies
                                                      across the membrane; THIRTEEN
                                                       predicted TM helices at aa 21-41
                                                      (NOT predicted using HMM to be a signal peptide), 66-88, 95-112, 127-149, 156-178, 188-205,
                                                       363-385, 389-411, 418-440,
```

444-462, 469-491, 501-523,

536-558; contains match to PROSITE

CDS

trypsin/alpha-amylase inhibitors family signature; some similarity to several transporters, e.g. Q9VG39, CG12286 protein (571 aa, Drosophila melanogaster, EMBL: AE003696, AAF54851); Fasta scores: E():1.8e-08, 27.1% identity in 188 aa' /codon-start=1 /label=L6071.01 /product="possible monocarboxylate transporter protein"
/protein-id="CAC32260.1"
/db-xref="GI:13122224"
/translation="MQIYEACKKADRAVTHRPAD **HWIGYLVAVSGALMQMMSYGIDNS** FSIFSNSMQNDPSLGYPSATTVSFGNSVSLGLSP **VFGILAGFLVDRVPPRVMMFTSTV** MLFAALWLSSSFAKSSAEVTASYSLLASISSALM LSPGAAATGSWFRRRLGLGOGINF CGGGVGSAVVPAVLGSLVDVYGWRHTFRLMSAFC AIGLVATILSCRRHPIEDDVDVDD HARGNNSPAREPSPDDCTAHRSPSHEERNEMMRM **ITSEAGENAAASPTTRMIDSMRTE** AEKAANRNSGDTITKGGKPDAAASARAALASPDG SDDMSMLLGRHQQQPQQLAQYGRK VHGTEACTVADLIQDMHARRLTWGEMMRVFLSVR **FLTHFFMFAIYGWSFYGLIYVAVP** YVSSMGSAGTVYAGVTPISTSKASTVFTFWGVFQ **IVGSILVGGVASFTDDALAYTMCA** TVGGLATSLLVFCRSYAAFAVCLSVVGFCTAGIF **AMMPALIAKDFHGPNLGFVMGCVF** VAGCLGGFSAPPIQAQLQTRYNGNYSYGCVFISC CTTFPGVLCYLLLWPAKQTRVGRV FTRVVRQA complement(1626..1691) /gene="L6071.01' /note="predicted TM helix region, aa 536-558' /note="region of BLASTN similarity to: AA756991 TENU0054 T.cruzi epimastigote normalized cDNA Library Trypanosoma cruzi cDNA clone 1k14 3', bases 3..306, 64% identity over 303 bases' complement(1731..1796) /gene="L6071.01' /note="predicted TM helix region, aa 501-523 /gene="L6071.01" /note="predicted TM helix region, aa 469-491" complement(1827..1892) /gene="L6071.01" complement(1884..1955) /note="PROSITE PS00426 Cereal trypsin/alpha-amylase inhibitors family signature complement(1914..1967) /gene="L6071.01" /note="predicted TM helix region, aa 444-462 /gene="L6071.01" complement(1980..2045) /note="predicted TM helix region, aa 418-440" complement(2067..2132) /gene="L6071.01" /note="predicted TM helix region, aa 389-411" complement(2145..2210) /gene="L6071.01" /note="predicted TM helix region, aa 363-385' /note="region of BLASTN similarity to: AQ847425 LMAJFV1-lm39e07.x1 Leishmania major FV1 random genomic library Leishmania major genomic clone LMAJFV1-lm39e07 3 similar to contains 3..462 V1-ch1-type-II.5 leishmania repetitive element;, bases 2..220, 98% identity over 218

bases region of BLASTN similarity

misc-feature

1666..1969

2189..2407

PS00426 Cereal

		to: AQ850931 LMAJFV1-lm32d03.x1
		Leishmania major FV1 random genomic library Leishmania major
	,	genomic clone LMAJFV1-lm32d03 3', bases 1385, 100% identity over
miss faatuun	complement(2685 2725)	384 bases" /gene="L6071.01"
misc-feature	complement(26852735)	/note="predicted TM helix region, aa 188-205"
misc-feature	complement(27662831)	/gene="L6071.01" /note="predicted TM helix region,
misc-feature	complement(28123185)	aa 156-178" /gene="L6071.01"
		/note="region of BLASTN similarity to: AW330418 TENU5164 T.cruzi
		epimastigote normalized cDNA
		Library Trypanosoma cruzi cDNA clone 9119 5', bases 1374, 67% identity over 373 bases region of
		BLASTN similarity to: AQ847139
		LMAJFV1-lm32d03.y1 Leishmania major FV1 random genomic library
		Leishmania major genomic clone
		LMAJFV1-1m32d03 5 similar to TR:Q08268 Q08268 CHROMOSOME XV
	÷ .	READING FRAME ORF YOL119C. ;, bases 1526, 98% identity over
		525 bases"
misc-feature	complement(28532918)	/gene="L6071.01" /note="predicted TM helix region,
misc-feature	complement(29643014)	aa 127-149" /gene="L6071.01"
misc-reacure	comprement(29043014)	/note="predicted TM helix region,
misc-feature	complement(30363101)	aa 95-112" /gene="L6071.01"
	-	/note="predicted TM helix region, aa 66-88"
misc-feature	complement(31773236)	/gene="L6071.01"
		/note="predicted TM helix region, aa 21-41"
repeat-region repeat-region	complement(33093330) complement(36013630)	<pre>/note="poly-pyrimidine tract" /note="poly-pyrimidine tract"</pre>
repeat-region	complement(37573774)	/note="poly-pyrimidine tract"
repeat-region misc-feature	complement(38933909) 40334247	<pre>/note="poly-pyrimidine tract" /note="region of BLASTN similarity</pre>
		to: AQ848Ž15 LMAJFV1-lm59c12.x1 Leishmania major FV1 random
		genomic library Leishmania major
		genomic clone LMAJFV1-lm59c12 3', bases 1215, 100% identity over
repeat-region	complement(41044121)	214 bases" /note="(acc)6"
repeat-region	complement(42084222)	/note="poly-pyrimidine tract"
misc-feature	complement(45164821)	/note="region of BLASTN similarity to: AQ851980 LMAJFV1-lm59c12.y1
		Leishmania major FV1 random genomic library Leishmania major
		genomic clone LMAJFV1-lm59c12 5',
	•	bases 1306, 99% identity over 305 bases"
repeat-region repeat-region	55355561 complement(55355552)	<pre>/note="poly-pyrimidine tract" /note="(ggggag)3"</pre>
repeat-region	56625681	/note="poly-pyrimidine tract"
repeat-region repeat-region	complement(56865697) complement(59105932)	<pre>/note="(atgt)3" /note="poly-pyrimidine tract" /note="poly-pyrimidine tract"</pre>
repeat-region gene	complement(64826496) complement(66518534)	/note="poly-pyrimidine tract" /gene="L6071.02"
CDS	complement(66518534)	/gene="L6071.02"
		/note="L6071.02, len = 626 aa, GTP-binding protein; contains Pfam
	•	match to entry PF01926 MMR-HSR1, GTPase of unknown function;
	•	contains match to PROSITE PS00017
		ATP/GTP-binding site motif A (P-loop); good similarity to many
		GTP-binding proteins, e.g.

NGP1-HUMAN, autoantigen ngp-1 (731 aa, Homo sapiens, EMBL: LÖ5425, AAC37588); Fasta scores: E():0, 43.5% identity in 646 aa' /codon-start=1 /label=L6071.02 /product="MMR-HSR1 GTP-binding protein" ./protein-id="CAC32261.1" /db-xref="GI:13122225" translation="MGKPGKKAGKGLLAPTNPNR/ RTDPNKTSLRDQRTIKRLKMYKSK **IKRDEKGNIIKGSVLKASDRIEQQMARIAPDRRW FGNTRTIGQEALQKFREEMGTKYK** DPYSVIIKQSKLPLSLLEEPKNTDGSIRKEMEWD KTFGDKANRKRVRLNAVDMSTLAT **EANVKGDYYDCNKKEKDRDLMKGVHKDRDDKTRN** GILMTKGQSNRIWCELYKVIDSSD **VVLYVVDARDPMGTRSAFLEDFMRREKKYKHFVL** VLNKCDLVPLWATARWLQILSKDY PTIAFHASVNHPFGKGNVISLLRQFARLHNVTHR **GSKRTKTPISVGVIGYPNVGKSSL** INTLRRKSVCKVAPIPGETKVWQYVALTRSIFLI DCPGVVYDRESNNDIQAVLKGVVR VERLGNADKTDVVDTVLKIVKQRDIVATYGVREW RDVVDFLEKLAKLRGKLVAGGEPD **VEAAARMLLYDWQRGRLPWFNAPPFESNKHHRDA** MEQPQEKHMKLIEHYSTFNVVDDT INRGDEKQDEGGDGDEETANNAADEDQLDSGSEA **EKDEEAVKPLKPSKTDRLSATKAD** TQLATVATYMREQEEKQRKAQRQQKRKAARKGQE DVEAFSADADRESDDALWAQFLAA AKV" /note="region of BLASTN similarity to: AL474947 TA177A02Q Trypanosoma brucei TREU927 sheared genomic DNA Trypanosoma brucei genomic clone 177a02 reverse, bases 236..576, 65% identity over 340 bases region of BLASTN similarity to: AQ660089 Sheared DNA-15K3.TR Sheared DNA Trypanosoma brucei genomic clone Sheared DNA-15K3, bases 139..535, 67% identity over 396 bases" /gene="L6071.02" /note="region of BLASTN similarity to: AQ660115 Sheared DNA-10N20.TF Sheared DNA Trypanosoma brucei genomic clone Sheared DNA-10N20, bases 1..233, 67% identity over 232 bases region of BLASTN similarity to: AQ645078 RPCI93-DpnII-29J3.TJ RPCI93-DpnII Trypanosoma brucei genomic clone RPCI93-DpnII-29J3, bases 1.384, 67% identity over 383 bases region of BLASTN similarity to: AQ638738 927P1-19F2.TP 927P1 Trypanosoma brucei genomic clone 927P1-19F2, bases 2..323, 70% identity over 321 bases region of BLASTN similarity to: BF936844 EST459899 Schistosoma mansoni female, Phil LoVerde/Joe Merrick Schistosoma mansoni cDNA clone SMFCC28 5' end, bases 442..611, 67% identity over 169 bases region of BLASTN similarity to: AI034991 TENG0051 T. cruzi epimastigote normalised cDNA Library Trypanosoma cruzi cDNA clone n33.r 5', bases 3..582, 70% identity over 579 bases" /gene="L6071.02" /note="Pfam match to entry PF01926 MMR-HSR1, GTPase of unknown function, score 272.20, E-value

misc-feature

6993..7333

misc-feature

complement(6993..7225)

misc-feature

complement(7116..7919)

6.8e-78'

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misc-feature
                     complement(7539...7562)
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                                                    /note="PROSITE PS00017
                                                    ATP/GTP-binding site motif A
                                                    (P-loop)"
misc-feature
                     complement(7619..7934)
                                                    /gene="L6071.02"
                                                    /note="region of BLASTN similarity
                                                    to: AI975898 EST270492 Schistosoma
                                                    mansoni female, Phil LoVerde/Joe
                                                    Merrick Schistosoma mansoni cDNA
                                                    clone SMFAY31 5' end, bases
                                                    109..424, 60% identity over 315
                                                    bases region of BLASTN similarity
                                                    to: BF936844 EST459899 Schistosoma
                                                    mansoni female, Phil Loverde/Joe
                                                    Merrick Schistosoma mansoni cDNA
                                                    clone SMFCC28 5' end, bases
                                                    109..424, 60% identity over 315
                                                    bases region of BLASTN similarity
                                                    to: AQ646760 RPCI93-ECORI-3H22.TJ
RPCI93-EcoRI Trypanosoma brucei
                                                    genomic clone RPCI93-EcoRI-3H22,
                                                    bases 45..354, 76% identity over 309 bases region of BLASTN
                                                    similarity to: AQ641764
                                                    RPCI93-ECORI-6J23.TJ RPCI93-ECORI
                                                    Trypanosoma brucei genomic clone RPCI93-EcoRI-6J23, bases 69..710, 72% identity over 641 bases region
                                                   of BLASTN similarity to: AQ647561
RPCI93-EcoRI-6I10.TJ RPCI93-EcoRI
Trypanosoma brucei genomic clone
RPCI93-EcoRI-6I10, bases 37..515,
70% identity over 478 bases region
                                                    of BLASTN similarity to: AQ641264
                                                    RPCI93-ECORI-5018.TJ RPCI93-ECORI
                                                    Trypanosoma brucei genomic clone
                                                    RPCI93-EcoRI-5018, bases 40..492, 70% identity over 452 bases region
                                                    of BLASTN similarity to: AQ642488
                                                    RPCI93-ECORI-3A8.TP RPCI93-ECORI
                                                   Trypanosoma brucei genomic clone RPCI93-EcoRI-3A8, bases 75..449, 71% identity over 374 bases"
misc-feature
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                                                    /note="region of BLASTN similarity
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                                                   Sheared DNA Trypanosoma brucei
                                                    genomic clone Sheared DNA-15K3,
                                                    bases 2..609, 65% identity over
                                                    607 bases'
misc-feature
                                                   /note="region of BLASTN similarity
to: AL161067 Leishmania major
                    7891..8389
                                                   Friedlin genomic clone cosmid
                                                   L4370 t3Hyg similar to
                                                   SW:NGP1-HUMAN Q13823 AUTOANTIGEN
                                                   NGP-1. [0] -1..., N=182,
                                                   Prob=1.9e-31, bases 14..512, 99%
                                                   identity over 498 bases"
/note="region of BLASTN similarity
misc-feature
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                                                   to: AQ942470 Sheared DNA-4515.TF
                                                   Sheared DNA Trypanosoma brucei
genomic clone Sheared DNA-4515,
                                                   bases 22..469, 68% identity over 447 bases"
                    9343..9358
repeat-region
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                    complement(9427..10017)
gene
                                                   /gene="L6071.03"
CDS
                    complement(9427..10017) /gene="L6071.03"
                                                   /note="L6071.03, len = 195 aa,
                                                   unknown; some similarity to
                                                   Q9SL93, putative kinetechore (163
                                                   aa, Arabidopsis thaliana, EMBL: AC006053, AAD31370); Fasta scores: E():0.57, 22.6% identity in 164
                                                   aa'
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/label=L6071.03

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                           KVDRDLYIIVHCDDGKYIEVSKNY
                           IKQCPFIEEAEGEIPEFGYPAAVLEHLIRWAVHY
                           GVDGHAASQLTRPCIYRDFSYVVT
                           DKWDNDFFNQRLCSPLNQKHYLLTMTAAEQFGMQ
                           GLLDFMCIGLGCKLRGKDDNGIIH
                           EVMGLDKEMEITSEDLAEVSRDYPWFDDAVKATT
                           KK"
complement(9861..10122)
                           /note="region of BLASTN similarity
                           to: AI034774 LmLv39p10/544D
                           Leishmania major promastigote full
                           length cDNA library from
                           stationary stage (day 10)
                           Leishmania major cDNA clone 544D 5', bases 1..262, 97% identity
                           over 261 bases region of BLASTN
                           similarity to: AIO34917
                           LmLv39p10/799C Leishmania major
                           promastigote full length cDNA
                           library from stationary stage (day
                           10) Leishmania major cDNA clone
                           799C 5', bases 1..262, 99%
                           identity over 261 bases region of
                           BLASTN similarity to: AI034649
                           LmLv39p10/369C Leishmania major
promastigote full length cDNA
                           library from stationary stage (day
                           10) Leishmania major cDNA clone
                           369C 5', bases 1..262, 99%
                           identity over 261 bases region of
                           BLASTN similarity to: AI034853
                           LmLv39p10/700D Leishmania major
                           promastigote full length cDNA
library from stationary stage (day
                           10) Leishmania major cDNA clone
                           700D 5', bases 1..250, 99%
                           identity over 249 bases region of
                           BLASTN similarity to: T93356
                           lmEST0116 LmLV39cDNA Leishmania
                           major cDNA clone Lm069 5' END,
                           bases 1..250, 99% identity over 249 bases"
complement(10499..13051 /gene="L6071.04"
complement(<10499..1305 /gene="L6071.04"
                           /note="L6071.04, len > 850 aa,
                           unknown; some similarity at amino
                           terminus to Q9RODO, smoothelin
                           large isoform 12 (921 aa, Mus
                           musculus, EMBL: AF132449,
AAF25578); Fasta scores:
E():0.041, 26.2% identity in 229
aa"
                           /codon-start=1
                           /label=L6071.04
                           /product="hypothetical protein
                           L6071.04"
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/db-xref="GI:13122227"
/translation="MSSTWLHALRAYAARTNATP
                           ASLAARCRLFDQLCAVQLRGLPSA
                           QLHPDALPVLREARWSLLREGTTADKAELLEHIV
                           RQYAQASEKAMYAAPTASCSLDGI
                           RGSADAQRQQPGKTSNHTRSLQDRSPQHATRSGT
                           SGSLLSGVQLSEMSVLQQLLLEQV
                           QCLVEAKVSDIPLSARHYRYPMRSPLLATVSPEL
                           PLLLLEELTQQSTAVSSSSSSSSS
                           AQSNIDWEARVDCCVGLVAAGHVQEALALCSDDG
```

STFRTVMHRVVPLRRDGWRCAWAL

MRHRAAVGARQVSDSANAERNHVF

ADAAPLSRVLDDATAAAGGTASLHWLRGVLEAVD

misc-feature

gene

DS

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LIMDTYLSVCPASRWRDAVGAVLE
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                                           SRSRNDSSPKQPDSDAPARWQVSDKSVQAKVEEC
                                           VRAARDADRLTVATGTILNERDKR
                                           HAAIYNHAMVALAATGHHTEAMHFYRTLPILLVN
                                           CYTHWSVLQLFLQPTRDSRAVSAL
                                           RSSENYNHCARALRHLIRMSTADSKAAHAHTQVN
                                           GNNGSVHPATPIRCTRDQGGVWES
                                           MILWAALRRDTETVDLCATHAPAVSRYAHLIALL
                                           SAAASRGDGWSAAQAQVRHMCAAP
                                           RTTLKELSLATAAMASFFPRWPAGTTAAELPVRA
                                           ELFDEVARSMAPLVGRSQSRMDEM
                                           LELLVGYSVSLRRRRRMPLTPQDEAAALDDILVK
                                           ENILANTMDLARPRAGYASDVDRA
                                           HHSCDNGAADDSGAWRTVVHVMTSVAERQGLSAA
                                           RAAPALVSAGVPAEMAIDL
misc-feature
                 11511..11979
                                           /note="region of BLASTN similarity
                                           to: AQ847561 LMAJFV1-lm40d09.x1
                                           Leishmania major FV1 random
                                           genomic library Leishmania major
                                           genomic clone LMAJFV1-lm40d09 3'
                                           bases 1..469, 99% identity over
                                           468 bases"
repeat-region
                 complement(11917..11928 /note="(tgtc)3"
repeat-region
                 complement(12402..12425 /note="(gtc)8"
misc-feature
                 complement(12792..13188 /note="region of BLASTN similarity
                                           to: AQ851423 LMAJFV1-lm40d09.y1
                                           Leishmania major FV1 random
                                           genomic library Leishmania major
                                           genomic clone LMAJFV1-lm40d09 5
                                           similar to contains Alu repetitive
                                           element;contains element
V1-ch1-type-II.7 leishmania
                                           repetitive element, bases 1..397, 100% identity over 396 bases"
repeat-region
                 complement(13081..13121 /note="poly-pyrimidine tract"
                 complement(13180..13195 /note="poly-pyrimidine tract"
repeat-region
gene
                 complement(13498..15009 /gene="L6071.05"
CDS
                 complement(13498..15009 /gene="L6071.05"
                                           /note="L6071.05, len = 502 aa,
                                           unknown; some similarity at amino
                                           terminus to Q9KY68, putative
                                           nlp/p60 family secreted protein
                                           (398 aa, Streptomyces coelicolor,
                                           ÈMBL: AĹ356832, CÁB92659); Fasta
                                           scores: E():2.5, 24.8% identity in
                                           141 aa'
                                           /codon-start=1
                                           /label=L6071.05
                                          /product="hypothetical protein
L6071.05"
                                          /protein-id="CAC32264.1"
/db-xref="GI:13122228"
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                                          YRDFLLEQRQREESQRQGGAGHAASSPCGGTSSD
                                          SSGDTLRADAVSFLQWALQYKLGG
                                          PNAFAHRRQELREMAAMMTEGCGAVTAAPPAAGA
                                          EAMRDGIAYDEDAVVAASFFTAAQ
                                          PGGVYYEYIRVGVLSVVLDGILFVHGGVNTSNAG
                                          FVPSLEATSYAEQVTAGQWWLPEV
                                          APQEVATTPTATSATGWLAALERFKAAAFSDWVN
                                          GAALRGEALRAYVYPRFVAPHSIA
                                          VGTVMNVDGPHYIPLTVVAYLLQSGIHTVCGGHQ
                                          PVGDTPAIIRQPGGFTIIDADNSY
                                          CGRGNKFCTRFNRRGAAVMELLFEHPDDHGGDEN
                                          VAPHDAVAAPSLTVHGYRADGAPF
```

EWVDRLRRLIVPSSKVALAGAEGQQQRVSATALR

EFDAYSDWRVGRYVGDGWWVRLPPEATAATSSLS

```
ETRWATAAEVDAWLRQAAASGKATVPGELAPRHT
                                              KEELAEVLAHRLKTKVKRT"
misc-feature
                   complement(13577..14073 /gene="L6071.05"
                                              /note="region of BLASTN similarity
                                              to: AQ853Ž48 LMAJFV1-lm81a10.y1
                                              Leishmania major FV1 random
                                              genomic library Leishmania major
                                              genomic clone LMAJFV1-lm81a10 5'
                                             similar to TR:Q42702 Q42702
                                             GLUCOSE-1-PHOSPHATE
                                             ADENYLYLTRANSFERASE PRECURSOR ;,
                                             bases 1..497, 99% identity over
                                              496 bases"
                  complement(15224..15238 /note="(cac)5"
repeat-region
repeat-region
                  complement(15614..15637 /note="poly-pyrimidine tract"
repeat-region
                  complement(15802..15817 /note="poly-pyrimidine tract"
misc-feature
                  complement(15936..16303 /note="region of BLASTN similarity
                                             to: AQ850713 LMAJFV1-lm41h09.x1
                                             Leishmania major FV1 random
                                             genomic library Leishmania major
                                             genomic clone LMAJFV1-lm41h09 3'
                                             bases 1..368, 96% identity over
                                             367 bases
misc-feature
                  16087..16257
                                             /note="region of BLASTN similarity
                                             to: AF008205 Leishmania major
                                             chromosome 1, complete sequence., bases 58478. 58648, 83% identity
                                             over 170 bases'
                  17246..17261
repeat-region
                                             /note="poly-pyrimidine tract"
                  complement(17631..17893 /note="region of BLASTN similarity
misc-feature
                                             to: AL161399 AL161399 Leishmania
                                             major Friedlin Leishmania major
                                             genomic clone PAC P719 right,
bases 1..263, 98% identity over
262 bases"
                  17967..17982 /note="poly-pyrimidine tract" complement(18162..18179 /note="poly-pyrimidine tract"
repeat-region
repeat-region
misc-feature
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                                             /note="region of BLASTN similarity
                                             to: AL455441 TA40D07Q Trypanosoma
                                             brucei TREU927 sheared genomic DNA
                                             Trypanosoma brucei genomic clone
                                             40d07 reverse, bases 144..462, 64% identity over 318 bases region of
                                             BLASTN similarity to: AL161165
                                             AL161165 Leishmania major Friedlin
                                             Leishmania major genomic clone cosmid L6812.2 t3Hyga similar to
                                             AP000373 AP000373 Arabidopsis
                                             thaliana genomic DNA,..., N=245
                                             Prob=3.1e-12, bases 1..422, 100%
                                             identity over 421 bases"
gene
                  complement(18229..18564 /gene="L6071.06"
CDS
                  complement(18229..18564 /gene="L6071.06"
                                             /note="L6071.06, len = 110 aa.
                                             possibly cytoplasmic dynein light
                                             chain; good similarity to many,
                                             e.g. Q9Ž336, tctex-1 (113 aa.
                                             Rattus norvegicus, EMBL: AB010119.
                                             BAA34532); Fāsta scores:
                                             E():1.2e-12, 35.5\% identity in 110
                                             aa'
                                             /codon-start=1
                                             /label=L6071.06
                                            /product="possible cytoplasmic dynein_light_chain"
                                             /protein-id="CAC32265.1"
                                             /db-xref="GI:13122229"
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FSGSEAAHAGLYELRRTODGFRHE

translation="MASGDRITLVDDASVICEDV"

```
DQVVQRLTQEAKLPRKYVVLVTILQKNGAGVQTI
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                                           IITVYGVTV"
repeat-region
                 complement(18630..18641 /note="(cca)4"
repeat-region
                 complement(18661..18678 /note="(cggcac)3"
repeat-region
                 complement(18679..18696 /note="(tggcac)3"
repeat-region
                 complement(18862..18886 /note="poly-pyrimidine tract"
                 complement(19320..20297 /gene="L6071.07"
gene
CDS
                 complement(19320..20297 /gene="L6071.07"
                                           /note="L6071.07, len = 324 aa,
                                           CDC27/NUC2-related protein;
                                           contains SIX Pfam matches to entry
                                          PF00515 TPR, TPR Domain; contains match to PROSITE PS00132 Zinc
                                           carboxypeptidases, zinc-binding
                                           region 1 signature; reasonable
                                           similarity to many, e.g.
                                           BIMA-EMENI, bima protein (806 aa
                                           Emericella nidulans, EMBL: X59269,
                                          CAA41959); Fasta scores:
E():2e-25, 31.0% identity in 290
                                           aa
                                           /codon-start=1
                                           /label=L6071.07
                                           /product="CDC27/NUC2-related
                                          protein'
                                           /protein-id="CAC32266.1"
                                           /db-xref="GI:13122230"
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                                          ATAGAVASTTSPWLLRQLALAHFH
                                          NGDIQESADAFEQLLRTAPWELTNPALIFYSTAL
                                          WHLKSESALGSLAQRLTDAEPLSA
                                          TTLCVVANAYSLIKDPRDALVMLKRAVQVAPTLA
                                          YAHALHGYELLGQDSKAEAEAEFK
                                          AALAVDASLYIAYAGLGERFMREEQIDKARGYYK
                                          EAVKLNPTPAIVNRFALTYHRQGK
                                          SLADLKTALRLYTESLERHPNNVTARRQRADVLL
                                          RLDQPMQALEELKALLVQCPGEAV
                                          VYVTLAECMVCLRRPHEALQHYQTAMHLDPRRES
                                          YVQGCIDQLVAANML'
misc-feature
                 complement(19374..19475 /gene="L6071.07"
                                           /note="Pfam match to entry PF00515
                                          TPR, TPR Domain, score 13.50,
                                          E-value 1.6"
misc-feature
                 19537..20187
                                          /note="region of BLASTN similarity
                                          to: AQ655661 Sheared DNA-2J6.TR
                                          Sheared DNA Trypanosoma, brucei
                                          genomic clone Sheared DNA-2J6,
                                          bases 27..677, 54% identity over
                                          650 bases
misc-feature
                 complement(19578..19670 /gene="L6071.07"
                                           /note="Pfam match to entry PF00515
                                          TPR, TPR Domain, score 2.20,
                                          E-value 26"
misc-feature
                 complement(19669..20187 /gene="L6071.07"
                                          /note="region of BLASTN similarity
                                          to: AQ658921 Sheared DNA-16B5.TR
                                          Sheared DNA Trypanosoma brucei
                                          genomic clone Sheared DNA-16B5,
                                          bases 138..656, 55% identity over 518 bases"
misc-feature
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                                          /note="Pfam match to entry PF00515
                                          TPR, TPR Domain, score 22.10,
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E-value 0.014'

VNALFSHETRYQHSKIAGLVSAIS

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misc-feature
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                                          E-value 31"
misc-feature
                complement(19833..19901 /gene="L6071.07"
                                          /note="PROSITE PS00132 Zinc
                                          carboxypeptidases, zinc-binding region 1 signature"
misc-feature
                complement(19893..19994 /gene="L6071.07"
                                          /note="Pfam match to entry PF00515
                                          TPR, TPR Domain, score 19.30, E-value 0.094"
                complement(20103..20204 /gene="L6071.07"
misc-feature
                                          /note="Pfam match to entry PF00515
                                          TPR, TPR Domain, score 13.00,
                                          E-value 1.8"
                complement(20271..20285 /note="poly-pyrimidine tract"
repeat-region
                 complement(20905..22083 /gene="L6071.08"
gene
                 complement(20905..22083 /gene="L6071.08"
CDS
                                          /note="L6071.08, len = 391 aa,
                                          unknown; some similarity to
                                          Q9UF25, hypothetical 22.4 Kd
                                          protein (222 aa, Homo sapiens,
                                          EMBL: AL133642, CAB63763); Fasta
                                          scores: E():0.082, 28.5% identity
                                          in 144 aa
                                          /codon-start=1
                                          /label=L6071.08
                                          /product="hypothetical protein
L6071.08"
                                          /protein-id="CAC32267.1
                                          /db-xref="GI:13122231"
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                                          HADPAASLAAAPPASHTSAPRATP
                                          DVATPPLPSASSAASPPLSSISSDVLSSLNASVQ
                                          ESLSGYLYTDAIELAQRLFDLEAS
                                          YAHLHLLAHCYTVSGATGTAYRLLQHYYPFLELH
                                          VTRPRTAGASTSAAGGAAGGVGSG
                                          VLPFDARRTWTASYGVHPSQHPFSSGMPSATAAV
                                          TFSASSNENNLTSDEFELGYETVD
                                          LQSQWDCQYLLGVCCYRTQHYEDGARVLSQLLYV
                                          CHQVTTTSSVLRRRLQQLRQQQQQ
                                          VTVRSDTSEEDDAGVAAAVRATEROLAGLCLRDG
                                          GAHFAGALLVGSVREASPAAPDRG
repeat-region
                 complement(21082..21096 /note="(cag)5"
repeat-region
                 complement(22052..22072 /note="(gcc)7"
misc-feature
                complement(22081..22191 /note="region of BLASTN similarity
                                          to: AQ846715 LMAJFV1-lm20b08.x1
                                          Leishmania major FV1 random
                                          genomic library Leishmania major
                                          genomic clone LMAJFV1-lm20b08 3'
                                          bases 1..111, 98% identity over
                                          110 bases
repeat-region
                complement(22223..22242 /note="poly-pyrimidine tract"
repeat-region
                 complement(22253..22267 /note="poly-pyrimidine tract"
repeat-region
                complement(23003..23018 /note="poly-pyrimidine tract"
gene
                complement(23178..24416 /gene="L6071.09"
CDS
                complement(23178..>2441 /gene="L6071.09"
                                          /note="L6071.09, len > 411 aa,
                                          possibly chromosome assembly
```

```
for N-terminal portion; predicted
                                             coiled-coil regions at aa 13-110,
                                             210-220; contains Pfam match to
                                             entry PF02483 SMC-C, SMC family
                                             C-terminal domain; contains match
to PROSITE PS00211 ABC
                                             transporters family signature;
                                             good similarity to several, e.g.
                                             BAB11491, chromosome assembly
                                             protein homolog (1175 aa,
                                             Arabidopsis thaliana, EMBL:
                                             AB019235, BAB11491); Fasta scores:
                                             E():2e-30, 35.4% identity in 378
                                             aa'
                                             /codon-start=1
                                             /label=L6071.09
                                             /product="chromosome segregation
                                             protein SMC2 homolog, C-terminal"
/protein-id="CAC32268.1"
/db-xref="GI:1312232"
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                                             DEERGAAEFERLEADMEQQAADLS
                                             RKTQDTEEDMVQQQSQKLKLAAQVEEVTQQLAAV
                                             QARSKQNEERRQRLEKDIDDAQEE
                                             LTRFAERKVTLDNLVKNGEVGLREQSRCLESLRR
                                             HVHEAEQRHSWLLEARATFNQPGG
                                             PYDFSDAARTAAILQELRDIEVRAAVMTSKLSQK
                                             SAILYEERRREYEELVKQRTALGE
                                             DKEAIQRCITEIESKKWGALDRMVGVVSSIFGKL
                                             FATCLPGATAQLLEERDAANHLSG
                                             LGVRVSFNGKPRESLSELSGGQRSLLALCLILAI
                                             LRVRPAPLYILDEVDAALDPSHTQ
                                             NIGRMLQLYFPHSQFLLVSLKDGMFNNANVLYHI
                                             RNTQGYSEVARIEHKPPPQPTSAD
                                             SDTRNVASGAENKDAVASFA'
misc-feature
                  complement(23271..23858 /gene="L6071.09"
                                             /note="Pfam match to entry PF02483
SMC-C, SMC family, C-terminal
                                             domain, score 88.10, E-value 1.8e-22"
misc-feature
                  complement(23493..23537 /gene="L6071.09"
                                             /note="PROSITE PS00211 ABC
                                             transporters family signature"
misc-feature
                  23503..23999
                                             /note="region of BLASTN similarity
                                             to: AL493037 TA342G06Q Trypanosoma
                                             brucei TREU927 sheared genomic DNA
                                             Trypanosoma brucei genomic clone
                  342g06 reverse, bases 2..498, 58% identity over 496 bases" complement(24517..24697 /note="region of BLASTN similarity
misc-feature
                                             to: AI034783 LmLv39p10/556D
                                             Leishmania major promastigote full
                                             length cDNA library from
                                             stationary stage (day 10)
                                             Leishmania major cDNA clone 556D
                                             5', bases 1..181, 86% identity
                                             over 180 bases region of BLASTN similarity to: AIO34933 LmLv39p10/815C Leishmania major
                                             promastigote full length cDNA
                                             library from stationary stage (day
                                             Leishmania major cDNA clone
                                             815C 5', bases 1..181, 94%
                                             identity over 180 bases"
                  complement(24735..24768 /note="poly-pyrimidine tract"
repeat-region
repeat-region
                  complement(25025..25053 /note="poly-pyrimidine tract"
repeat-region
                  complement(25041..25052 /note="(cct)4"
                  complement(25404..25415 /note="(agt)4"
repeat-region
```

complement(25416..25435 /note="poly-pyrimidine tract"

repeat-region

protein SMC2 homolog; see L6071.12

```
repeat-region
                  complement(25607..25626 /note="poly-pyrimidine tract"
gene
                  complement(25835..26614 /gene="L6071.10"
                  complement(25835..26614 /gene="L6071.10"
CDS
                                             /note="L6071.10, len = 258 aa,
                                             unknown; some similarity to
                                             OREX-RAT, orexin precursor (130
                                             aa, Rattus norvegicus, EMBL:
                                             AFÓ19565, AACO2933); Fasta scores:
E():1, 25.2% identity in 127 aa"
                                             /codon-start=1
                                             /label=L6071.10
                                             /product="hypothetical protein
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/db-xref="GI:13122233"
                                             translation="MPPAPAPPFSSVSQCLSALL"
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                                             HIIFSEPHIVFYCVCPFTRCHRRRPLPSPLSLSL
                                             PQHTTLSTAINKHEAEVVLDWSMS
                                             RLARALAKPFTVPVAMCTRHVAAMDEPLKRHIDA
                                             YAARGEDITIAVWREYVDGQRALL
                                             PYRWTKFRSEVAYLTSGQMAITDLTFADLLVFIR
                                             FLTKCLFIFIVAVMVGRRSVFPSL
                                             EPTSPFVEEIVKNWQPNRLHGVAGAEYMACDQAA
                                             AAGYGHR"
epeat-region
                 complement(26377..26411 /note="poly-pyrimidine tract"
                 complement(26823..26841 /note="poly-pyrimidine tract"
epeat-region
repeat-region
                 complement(27165..27179 /note="poly-pyrimidine tract"
repeat-region
                 complement(27484..27505 /note="poly-pyrimidine tract"
repeat-region
                 complement(27627..27641 /note="poly-pyrimidine tract"
epeat-region
                 complement(28094..28105 /note="(caaq)3"
epeat-region
                 complement(28155..28166 /note="(gtat)3"
nisc-feature
                 28343..28575
                                             /note="region of BLASTN similarity
                                             to: AQ849379 LMAJFV1-lm47b03.y1
                                             Leishmania major FV1 random
                                             genomic library Leishmania major
                                             genomic clone LMAJFV1-lm47b03 5
                                             bases 160..392, 100% identity over
                                             232 bases"
                 28443..28463 /note="poly-pyrimidine tract"
29010..29031 /note="poly-pyrimidine tract"
complement(29105..29459 /note="region of BLASTN similarity
repeat-region
repeat-region
isc-feature
                                             to: AQ902240 LMAJFV1-lm47b03.x1
                                             Leishmania major FV1 random
                                             genomic library Leishmania major
                                             genomic clone LMAJFV1-lm47b03~3'
                                             similar to contains element
                                             V1-ch1-type-II.14 leishmania
                                             repetitive element, bases 1..355, 100% identity over 354 bases
                                             region of BLASTN similarity to:
                                             AQ940065 Sheared DNA-19A7.TF
                                            Sheared DNA Trypanosoma brucei genomic clone Sheared DNA-19A7,
                                             bases 1..284, 79% identity over
                                            283 bases region of BLASTN
                                             similarity to: AL472284 TA161B10P
                                             Trypanosoma brucei TREU927 sheared
                                             genomic DNA Trypanosoma brucei
                                             genomic clone 161b10 forward,
                                            bases 39..398, 78% identity over 359 bases region of BLASTN
                                             similarity to: AL463943 TA116C11Q
                                            Trypanosoma brucei TREU927 sheared
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genomic DNA Trypanosoma brucei

bases 237..522, 78% identity over 285 bases" /note="poly-pyrimidine tract"
/gene="L6071.11"
/gene="L6071.11"
/note="L6071.11, len = 573 aa, 29113..29129 29147..30871 29147..30871 repeat-region gene CDS ATPase alpha subunit; contains Pfam match to entry PF00006 ATP-synt-ab, ATP synthase alpha/beta family, Pfam match to entry PF00422 ATP-synt-A-c, ATP synthase Alpha chain, C terminal; contains match to PROSITE PS00152 ATP synthase alpha and beta subunits signature, PROSITE PS00070 Aldehyde dehydrogenases cysteine active site, PROSITE PS00017 ATP/GTP-binding site motif A (P-loop); good similarity to many, e.g. AAG23339, ATPase alpha subunit (585 aa, Trypanosoma brucei brucei, EMBL: AY007705 AAG23339); Fasta scores: E():0, 88.6% identity in 587 aa' /codon-start=1 /label=L6071.11 /product="ATPase alpha subunit" /protein-id="CAC32270.1" /db-xref="GI:13122234" /translation="MRRFVAQYVAPAMGRLASTA **AAGKSAAPGQKSFFKATEMIGYVH** SIDGTIATLIPAPGNPGVAYNTIIMIQVSPTTFA AGLVFNLEKDGRIGIILMDNITEV QSGQKVMATGKLLYIPVGAGVLGKVVNPLGHEVP VGLLTRSRALLESEQTLGKVDAGA PNIVSRSPVNYNLLTGFKAVDTMIPIGRGQRELI VGDRQTGKTSIAVSTIINQVRSNQ QILSKNAVISIYVSIGQRCSNVARIHRLLRSYGA LRYTTVMAATAAEPAGLQYLAPYS GVTMGEYFMNRGRHCLCVYDDLSKQAVAYRQISL LLRRPPGREAYPGDVFYLHSRLLE RAAMLSPGKGGGSVTALPIVETLSNDVTAYIVTN VISITDGQIYLDTKLFTGGQRPAV NIGLSVSRVGSSAQNVAMKAVAGKLKGILAEYRK LAADSVGGSQVQTVPMIRGARFVA LFNQKNPSFFMNALVSLYACLNGYLDDVKVNYAK LYEYLLVNKDLSVMYGTATNKFFY MYVQQLNYVIRFFTLNHPILNAEVEEMLKQHTHL FLQHYQSKMNAIKTEKEIKALKNL LYSCKRAV" /gene="L6071.11" nisc-feature 29234...29420 /note="region of BLASTN similarity to: AL472354 TA161B10Q Trypanosoma brucei TREU927 sheared genomic DNA Trypanosoma brucei genomic clone 161b10 reverse, bases 327..513, 79% identity over 186 bases region of BLASTN similarity to: AQ658777 Sheared DNA-27D17.TF Sheared DNA Trypanosoma brucei genomic clone sheared DNA-27D17, bases 91..450, 78% identity over 359 bases region of BLASTN similarity to: AY007705 Trypanosoma brucei brucei ATPase alpha subunit mRNA, complete cds; nuclear gene for kinetoplast product., bases 124..483, 79% identity over 359 bases' /gene="L6071.11" nisc-feature 29291..30346 /note="Pfam match to entry PF00006 ATP-synt-ab, ATP synthase alpha/beta family, score 420.30, E-value 1.8e-122" nisc-feature complement(29600..30196 /note="region of BLASTN similarity to: AQ943100 Sheared DNA-35K24.TR

genomic clone 116c11 reverse,

Sheared DNA Trypanosoma brucei

bases 1..597, 79% identity over 596 bases" complement(29600..29826 /note="region of BLASTN similarity to: AL463943 TA116C110 Trypanosoma brucei TREU927 sheared genomic DNA misc-feature Trypanosoma brucei genomic clone 116c11 reverse, bases 10..236, 79% identity over 226 bases" misc-feature 29664..30267 /gene="L6071.11" /note="region of BLASTN similarity to: AI976759 EST271353 Schistosomá mansoni male, Phil LoVerde/Joe Merrick Schistosoma mansoni cDNA clone SMMAA46 5' end similar to ATP synthase alpha subunit, bases 94..697, 57% identity over 603 bases region of BLASTN similarity to: AY007705 Trypanosoma brucei brucei ATPase alpha subunit mRNA, complete cds; nuclear gene for kinetoplast product., bases 619..1758, 80% identity over 1139 bases region of BLASTN similarity to: AA186203 T3871 MVAT4 bloodstream form of serodeme WRATat1.1 Trypanosoma brucei rhodesiense CDNA 5' similar to gi|45606 (X66103) ATPase alpha subunit [Propionigenium modestum], bases 1..297, 79% identity over 296 bases region of BLASTN similarity to: AQ950429 Sheared DNA-39I2.TF Sheared DNA Trypanosoma brucei genomic clone Sheared DNA-39I2, bases 1..410, 79% identity over 409 bases region of BLASTN similarity to: AZ050719 GSSTc11570 Trypanosoma cruzi random genomic library Trypanosoma cruzi genomic clone G38N18, bases 1..213, 83% identity over 212 bases" misc-feature complement(29665..30067 /note="region of BLASTN similarity to: AL161137 AL161137 Leishmania major Friedlin genomic clone cosmid L6071.2 t3Hyga similar to SLATPSYNA Z22606 S.lividans i protein and ATP synthase... N=824, Prob=3.3e-60; SW:ATPA-BOVIN P19482 ATP SYNTHASE ALPHA CHAIN LIVER..., N=424, Prob=1.3e-52, bases 1..403, 100% identity over 402 bases region of BLASTN similarity to: AQ950431 Sheared DNA-39I2.TR Sheared DNA Trypanosoma brucei genomic clone Sheared DNA-39I2, bases 2..483, 79% identity over 481 bases" /gene="L6071.11" misc-feature 29732..29755 /note="PROSITE PS00017 ATP/GTP-binding site motif A (P-loop) nisc-feature 29829..30350 /gene="L6071.11" /note="region of BLASTN similarity to: AQ653001 Sheared DNA-1M5.TF Sheared DNA Trypanosoma brucei genomic clone Sheared DNA-1M5, bases 1..522, 78% identity over 521 bases region of BLASTN similarity to: AA556054 TENF0235 T.cruzi epimastigote normalized cDNA Library Trypanosoma cruzi cDNA clone 235 5' similar to ATP synthase, bases 10..418, 78%

genomic clone Sheared DNA-35K24,

identity over 408 bases region of

BLASTN similarity to: N45888 T1405 MVAT4 bloodstream form of serodeme WRATat1.1 Trypanosoma brucei rhodesiense cDNA 5' similar to Na+-transporting ATP synthase alpha chain, bases 24..292, 77% identity over 268 bases region of BLASTN similarity to: AI067305 EST208983 Schistosoma mansoni, Phil LoVerde/Joe Merrick Schistosoma mansoni cDNA clone SMNAS35 5' end similar to ATP synthase, alpha subunit, bases 4..420, 59% identity over 416 bases region of BLASTN similarity to: AI067846 EST209530 Schistosoma mansoni, Phil LoVerde/Joe Merrick Schistosoma mansoni cDNA clone SMNCF75 5' end similar to ATP synthase, alpha subunit, bases 4. 457, 59% identity over 453 bases region of BLASTN similarity to: AI068328 EST210019 Schistosoma mansoni, Phil LoVerde/Joe Merrick Schistosoma mansoni cDNA clone SMNCT59 5' end similar to ATP synthase, alpha subunit, bases 4.457, 59% identity over 453 bases region of BLASTN similarity to: AI067947 EST209635 Schistosoma mansoni, Phil LoVerde/Joe Merrick Schistosoma mansoni cDNA clone SMNCH12 5' end similar to ATP synthase, alpha subunit, bases 4..499, 57% identity over 495 bases re gion of BLASTN similarity to: AI068269 EST209960 Schistosoma mansoni, Phil LoVerde/Joe Merrick Schistosoma mansoni cDNA clone SMNCQ92 5' end similar to ATP synthase, alpha subunit, bases 10..505, 58% identity over 495 bases region of BLASTN similarity to: AL160498 AL160498 Leishmania major Friedlin Leishmania major genomic clone PAC P108 right similar to MXPTATP D16176 M.xanthus DNA for proton translocating..., N=1252, Prob=6.5e-96; SW:ATPA-DROME P35381 ATP SYNTHASE ALPHA CHAIN,. N=548, Prob=4.3e-69, bases 3..450, 100% identity over 447 bases region of BLASTN similarity to: AZ215908 Sheared DNA-116C10.TF Sheared DNA Trypanosoma brucei genomic clone Sheared DNA-116c10, bases 2..633, 79% identity over 631 bases" /gene="L6071.11" /note="PROSITE PS00070 Aldehyde dehydrogenases cysteine active site' /gene="L6071.11" /note="region of BLASTN similarity to: AZ050667 GSSTc11516 Trypanosoma cruzi random genomic library Trypanosoma cruzi genomic clone G36G9, bases 1..336, 84% identity over 335 bases" complement(30200..30568 /note="region of BLASTN similarity to: AZ051030 GSSTc11882 Trypanosoma cruzi random genomic

library Trypanosoma cruzi genomic clone G54D16, bases 1..369, 84%

misc-feature

29834..29869

misc-feature

30156..30491

misc-feature

misc-feature	complement(3029030870)	identity over 368 bases" /note="region of BLASTN similarity to: AQ658781 Sheared DNA-27D17.TR Sheared DNA Trypanosoma brucei genomic clone Sheared DNA-27D17, bases 91671, 79% identity over 580 bases region of BLASTN similarity to: AZ218156 Sheared DNA-58H4.TR Sheared DNA Trypanosoma brucei genomic clone Sheared DNA-58H4, bases 164576, 78% identity over 412 bases region of BLASTN similarity to: AA875724 TENU0195 T.cruzi epimastigote normalized cDNA Library Trypanosoma cruzi cDNA clone 6e3 3', bases 197477, 81% identity over 280 bases region of BLASTN similarity to: AI021883 TENU0477 T. cruzi epimastigote normalized cDNA Library Trypanosoma cruzi cDNA clone 7e21 3', bases 190424, 79% identity over 234
	•	bases region of BLASTN similarity to: AQ911949 LMAJFV1-ln07g01.y1 Leishmania major FV1 random genomic library Leishmania major genomic clone LMAJFV1-ln07g01 5', bases 1239, 99% identity over
misc-feature	3031430343	238 bases" /gene="L6071.11" /note="PROSITE PS00152 ATP synthase alpha and beta subunits
misc-feature	3034730433	signature" /gene="L6071.11" /note="Pfam match to entry PF00422 ATP-synt-A-c, ATP synthase Alpha chain, C terminal, score 24.40,
misc-feature	3037330582	E-value 2.3e-07" /gene="L6071.11" /note="region of BLASTN similarity to: AQ940066 Sheared DNA-19A7.TR Sheared DNA Trypanosoma brucei genomic clone Sheared DNA-19A7, bases 1.210, 78% identity over
misc-feature	3042830748	209 bases" /gene="L6071.11" /note="region of BLASTN similarity to: AI073316 TENU2987 T. cruzi epimastigote normalized cDNA Library Trypanosoma cruzi cDNA clone 28c8 5', bases 1321, 80% identity over 320 bases region of BLASTN similarity to: AI035122 TENG0213 T. cruzi epimastigote normalised cDNA Library Trypanosoma cruzi cDNA clone n250.r 5', bases 1367, 80%
misc-feature	3083731319	identity over 366 bases' /note="region of BLASTN similarity to: AZ082002 L2005k.d-HygT7.1 Leishmania major Friedlin Cosmid Genomic Library Leishmania major genomic clone L2005k, bases 11493, 98% identity over 482 bases"
repeat-region repeat-region misc-feature		/note="(agt)4" /note="(acc)6" /note="region of BLASTN similarity to: AL491879 TA320H10Q Trypanosoma brucei TREU927 sheared genomic DNA Trypanosoma brucei genomic clone 320h10 reverse, bases 133536,
gene CDS	3182534155	76% identity over 403 bases" /gene="L6071.12" /gene="L6071.12"

/note="L6071.12, len > 776 aa, possibly chromosome segregation protein SMC2 homolog; see L6071.09 for C-terminal portion; predicted extensive multiple coiled-coil regions at aa 180-480, 711-776, continued in L6071.09; contains Pfam match to entry PF02463 SMC-N, SMC domain N terminal domain; contains match to PROSITE PSÓ1156 TonB-dependent receptor proteins signature 2, PROSITE PS00017 ATP/GTP-binding site motif A (P-loop); good similarity to many, e.g. SMCŹ-YEAST, chromosome segregation protein smc2 (1170 aa, Saccharomyces cerevisiae, EMBL: D44602, BÁA08042); Fasta scores: E():0, 29.2% identity in 811 aa" /codon-start=1 /label=L6071.12 /product="chromosome segregation protein SMC2 homolog, N-terminal" /protein-id="CAC32271.1" /db-xref="GI:13122235" /translation="MRVKSIVIDGFKSYAHRKEL ADLSPHFNAITGLNGSGKSNIFDA **ICFVMGITNLKRVRAEDPRELIFRAGTTGVHAAR** VTIEFVNDDPASAPPGYSCEEYPL ITIGRQIKLGGRQQFFFNNTVSLQSKVKRFFESI SLNVDNPHFMILQGTVHKLIGMRS QDILSLIEEAVGTKAFDHRRRTAETLIRNKERKM EEIDTNIEAQIRPLLETMRADQEE YNTFMQMREKMEEKVRFRVALDYHTHRTQHAEAE AAMTARKADVQNAKTQLQALPRQE EEAARRLLQLQDSLSAPSEAAIALHEEEDELKKA **HSRLEGQLGNCTKSLKQLETQLKS** LRKEQERQSSSQAAFAARQREHEQLLAQIKEGKE **TCAKLKKGLKLLRSGVQAGASGVS** LAEERQQVDLQLIEQQSRVRRATDRLEELVKQQR RVEAHQAEESSRVRHLEREYAKAT ASLEKAKAVYTPLALKQQRKEALEAEISSLKREC QAEYENFQRQVSTATARNYDLDYN RYACPPDTEDKVLGRVGQLITPTDPQHALGLMVG AQNQLLRVVVTDDRVAEAIIRSGL RQRTAFFALDKLQRQPTHFFIDGAKLQAARLMAE QQGGWVHRARDLVTVQEASSHQQQ QQLNALADFVFGNFLVCSSLRLAQELAYDASIKA KAVTVEGEVAEPNGLMTGGSTRQL RDVFADLKTYTAQKEPLKALQQRTRALEVEYAAL RDTLRQHQHDIQVYKTAEEAAELS KQRYIVAANSAQSGAAEQAEQIEREQTALAEARE KVE" /gene="L6071.12" /note="Pfam match to entry PF02463 SMC-N, SMC domain N terminal domain, score 176.70, E-value 3.9e-49" /gene="L6071.12" /note="PROSITE PS00017 ATP/GTP-binding site motif A (P-loop)"

complement(31950..32566 /note="region of BLASTN similarity to: AZ215185 Sheared DNA-83G9.TR Sheared DNA Trypanosoma brucei genomic clone Sheared DNA-83G9. bases 2..618, 71% identity over 616 bases region of BLASTN similarity to: AQ649957 Sheared DNA-5H11.TR Sheared DNA Trypanosoma brucei genomic clone Sheared DNA-5H11, bases 80..689, 61% identity over 609 bases region of BLASTN similarity to: AQ653Ö51 Sheared DNA-6M4.TR Sheared DNA

Trypanosoma brucei genomic clone

misc-feature

31825..32343

misc-feature

31918..31941

misc-feature

misc-feature 32122..32175

Sheared DNA-6M4, bases 27..679, 54% identity over 652 bases" /gene="L6071.12" /note="PROSITE PS01156 TonB-dependent receptor proteins signature 2" /note="(cag)5"

repeat-region 33688..33702

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    AUTHOR (AU):
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human

narcoleptic brains

TITLE (TI):

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Nat. Med., 6 (9), 991-997 ( ***2000*** )
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                           Department of Psychiatry, Stanford University Medical
                           Center, 1201 Welch Road, MSLS Bldg. P112, Stanford, CA
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DATE (DATE):
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                         A mutation in a case of early onset narcolepsy and a generalized absence of ***hypocretin*** peptides
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                              Peyron,C.; Faraco,J.; Rogers,W.; Ripley,B.; Overeem,S.;
   AUTHOR (AU):
                              Charnay,Y.; Nevsimalova,S.; Aldrich,M.; Reynolds,D.; Albin,R.; Li,R.; Hungs,M.; Pedrazzoli,M.; Padigaru,M.; Kucherlapati,M.; Fan,J.; Maki,R.; Lammers,G.J.;
                              Bouras, C.; Kucherlapati, R.; Nishino, S.; Mignot, E.
                              A mutation in a case of early onset narcolepsy and a neneralized absence of ***hypocretin*** peptides
   TITLE (TI):
                              generalized absence of 
***human*** narcole
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                                                   narcoleptic brains
                               Nat. Med., 6 (9), 991-997 (
                                                                   ***2000*** )
   JOURNAL (SO):
                              CA 133:348631
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   AUTHOR (AU):
                              Faraco, J.; Rogers, W.; Overeem, S.; Li, R.; Mignot, E.
   TITLE (TI):
                              Direct Submission
                              Submitted (05-NOV-1999) Center for Narcolepsy Research, Department of Psychiatry, Stanford University Medical Center, 1201 Welch Road, MSLS Bldg. P112, Stanford, CA
   JOURNAL (SO):
                               94305-5485, USA
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L5
      ANSWER 107 OF 154
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                                                   COPYRIGHT 2004 on STN
                               F202078S11
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DIVISION CODE (CI):
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DATE (DATE):
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DEFINITION (DEF):
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                               11 of 14
SEGMENT:
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SOURCE:
ORGANISM (ORGN):
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Bouras, C.; Kucherlapati, R.; Nishino, S.; Mignot, E.
A mutation in a case of early onset narcolepsy and a generalized absence of ***hypocretin*** peptides

***human*** narcoleptic brains
   TITLE (TI):
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   JOURNAL (SO):
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   TITLE (TI):
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    ANSWER 108 OF 154
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                           6 Feb 2001
DATE (DATE):
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DEFINITION (DEF):
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EGMENT:
                           10 of 14
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OURCE:
ORGANISM (ORGN):
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                          Hominidae; Homo
74 a 85 c
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NUCLEIC ACID COUNT (NA): 74 a
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                          Bouras, C.; Kucherlapati, R.; Nishino, S.; Mignot, E.
A mutation in a case of early onset narcolepsy and a
generalized absence of ***hypocretin*** peptides
  TITLE (TI):
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  TITLE (TI):
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  JOURNAL (SO):
                          Department of Psychiatry, Stanford University Medical Center, 1201 Welch Road, MSLS Bldg. P112, Stanford, CA 94305-5485, USA
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SEQUENCE LENGTH (SQL):
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MOLECULE TYPE (CI):
DIVISION CODE (CI):
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DATE (DATE):
                            6 Feb 2001
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DEFINITION (DEF):
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SEGMENT:
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REFERENCE:
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   JOURNAL (SO):
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                            Department of Psychiatry, Stanford University Medical
                            Center, 1201 Welch Road, MSLS Bldg. P112, Stanford, CA
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GenBank VERSION (VER): AF202085
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DIVISION CODE (CI):
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DEFINITION (DEF):
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SEGMENT:
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L5

SOURCE:

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                            Bouras, C.; Kucherlapati, R.; Nishino, S.; Mignot, E.
                            A mutation in a case of early onset narcolepsy and a generalized absence of ***hypocretin*** peptides ***human*** narcoleptic brains
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CA 133:348631
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    AUTHOR (AU):
                            Faraco, J.; Rogers, W.; Overeem, S.; Li, R.; Mignot, E.
    TITLE (TI):
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    JOURNAL (SO):
                            Submitted (05-NOV-1999) Center for Narcolepsy Research,
                            Department of Psychiatry, Stanford University Medical
                            Center, 1201 Welch Road, MSLS Bldg. P112, Stanford, CA
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LOCUS (LOC):
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DATE (DATE):
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SEGMENT:
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A mutation in a case of early onset narcolepsy and a generalized absence of ***hypocretin*** peptides
    TITLE (TI):
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    TITLE (TI):
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      ANSWER 112 OF 154
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GenBank ACC. NO. (GBN): AF202083
GenBank VERSION (VER):
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DATE (DATE):
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DEFINITION (DEF):
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                           6 of 14 ***human***
SEGMENT:
SOURCE:
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NUCLEIC ACID COUNT (NA): 45 a
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                           Albin,R.; Li,R.; Hungs,M.; Pedrazzoli,M.; Padigaru,M.; Kucherlapati,M.; Fan,J.; Maki,R.; Lammers,G.J.; Bouras,C.; Kucherlapati,R.; Nishino,S.; Mignot,E. A mutation in a case of early onset narcolepsy and a generalized absence of ***hypocretin*** peptides in
   TITLE (TI):
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   JOURNAL (SO):
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                           Submitted (05-NOV-1999) Center for Narcolepsy Research,
                           Department of Psychiatry, Stanford University Medical
Center, 1201 Welch Road, MSLS Bldg. P112, Stanford, CA
94305-5485, USA
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LOCUS (LOC):
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DIVISION CODE (CI):
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DATE (DATE):
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SEGMENT:
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Hominidae; Homo

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    AUTHOR (AU):
                              Bouras, C.; Kucherlapati, R.; Nishino, S.; Mignot, E.
                              A mutation in a case of early onset narcolepsy and a neneralized absence of ***hypocretin*** peptides
    TITLE (TI):
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                                 ***human***
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                              Nat. Med., 6 (9), 991-997 ( ***2000*** )
    JOURNAL (SO):
    OTHER SOURCE (OS):
                              CA 133:348631
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                              Faraco,J.; Rogers,W.; Overeem,S.; Li,R.; Mignot,E.
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    TITLE (TI):
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                              Submitted (05-NOV-1999) Center for Narcolepsy Research, Department of Psychiatry, Stanford University Medical
    JOURNAL (SO):
                              Center, 1201 Welch Road, MSLS Bldg. P112, Stanford, CA
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      ANSWER 114 OF 154
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DATE (DATE):
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                              4 of 14
***human***
SEGMENT:
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A mutation in a case of early onset narcolepsy and a
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                              Faraco, J.; Rogers, W.; Overeem, S.; Li, R.; Mignot, E.
                             Direct Submission
   JOURNAL (SO):
                             Submitted (05-NOV-1999) Center for Narcolepsy Research,
                             Department of Psychiatry, Stanford University Medical
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     ANSWER 115 OF 154
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LOCUS (LOC):
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SEQUENCE LENGTH (SQL):
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MOLECULE TYPE (CI):
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DIVISION CODE (CI):
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DATE (DATE):
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DEFINITION (DEF):
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                                                                    ***receptor*** -1
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                            (HCRTR1) gene, exon 3.
3 of 14
***human***
SEGMENT:
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NUCLEIC ACID COUNT (NA): 58 a
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                           Peyron, C.; Faraco, J.; Rogers, W.; Ripley, B.; Overeem, S.;
                           Charnay, Y.; Nevsimalova, S.; Aldrich, M.; Reynolds, D.; Albin, R.; Li, R.; Hungs, M.; Pedrazzoli, M.; Padigaru, M.; Kucherlapati, M.; Fan, J.; Maki, R.; Lammers, G.J.; Bouras, C.; Kucherlapati, R.; Nishino, S.; Mignot, E. A mutation in a case of early onset narcolepsy and a generalized absence of ***hypocretin*** peptides in
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   AUTHOR (AU):
   TITLE (TI):
   JOURNAL (SO):
                           Submitted (05-NOV-1999) Center for Narcolepsy Research,
                           Department of Psychiatry, Stanford University Medical
Center, 1201 Welch Road, MSLS Bldg. P112, Stanford, CA
                           94305-5485, USA
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     ANSWER 116 OF 154
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LOCUS (LOC):
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DATE (DATE):
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DEFINITION (DEF):
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SEGMENT:
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48 a 98 c
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  AUTHOR (AU):
                           Kucherlapati,M.; Fan,J.; Maki,R.; Lammers,G.J.
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A mutation in a case of early onset narcolepsy and a
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  TITLE (TI):
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  JOURNAL (SO):
                           Department of Psychiatry, Stanford University Medical
                           Center, 1201 Welch Road, MSLS Bldg. P112, Stanford, CA
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    ANSWER 117 OF 154
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OCUS (LOC):
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A mutation in a case of early onset narcolepsy and a
generalized absence of ***hypocretin*** peptides
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Nat. Med., 66 (9), 991-997 ( ***2000*** )
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   JOURNAL (SO):
                           Submitted (05-NOV-1999) Center for Narcolepsy Research,
                          Department of Psychiatry, Stanford University Medical
                           Center, 1201 Welch Road, MSLS Bldg. P112, Stanford, CA
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DATE (DATE):
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DEFINITION (DEF):
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SOURCE:
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                          Sakurai,T.; Amemiya,A.; Ishii,M.; Matsuzaki,I.; Chemelli,R.M.; Tanaka,H.; Williams,S.C.;
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                          Richardson,J.A.; Kozlowski,G.P.; Wilson,S.;
                          Arch, J.R.S.; Buckingham, R.E.; Haynes, A.C.; A.
                          Carr,S.A.; Annan,R.S.; McNulty,D.E.; Liu,W.-S.;
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                          Orexins and ***orexin***
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hypothalamic neuropeptides and G protein-coupled
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                                                  Direct Submission
      TITLE (TI):
                                                  Submitted (07-JAN-1998) HHMI/Department of Molecular Genetics, University of Texas Southwestern Medical Center at Dallas, 5323 Harry Hines Blvd., Rm. Y5.224,
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   TITLE (TI):
                             Orexins and
                                                                 receptors: a family of
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                             receptors that regulate feeding behavior Cell, 92 (4), 573-585 ( ***1998*** ) CA 128:290571
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   TITLE (TI):
                             Direct Submission
   JOURNAL (SO):
                             Submitted (07-JAN-1998) HHMI/Department of Molecular Genetics, University of Texas Southwestern Medical
                             Center at Dallas, 5323 Harry Hines Blvd., Rm. Y5.224,
                             Dallas, TX 75235-9050, USA
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         PHENYL UREA AND PHENYL THIOUREA DERIVATIVES; PHENYL UREA AND PHENYL
         THIOUREA DERIVATIVES AND THEIR USE AS PHARMACEUTICALS
         Coulton Steven (GB); Johns Amanda (GB); Porter Roderick Alan (GB)
 IN
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                              B1 20030722
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        PHENYLUREA AND PHENYLTHIO UREA DERIVATIVES:
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        US 6372757
                              B1 20020416
        WO 9958533
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     ASAHI S; EGASHIRA S; MATSUDA M; IWAASA H; KANATANI A; OHKUBO M; IHARA M;
ΑU
     MORISHIMA H
     SAKURAI T
CS
     Banyu Tsukuba Res. Inst., Ibaraki, Jpn
     Univ. Tsukuba, Ibaraki, Jpn
     Pept Sci, (2000) vol. 1999, pp. 37-40. Journal Code: X0695A (Fig. 1. Tbl.
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AN
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ΤI
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     neuropeptides that control feeding behavior.
AU 1
     SAKURAI TAKESHI
     Univ. of Tsukuba, Inst. of Basic Med. Sci. Saibo Kogaku (Cell Technology), (1998) vol. 17, no. 6, pp. 864-865.
CS
S0
     Journal Code: Y0229A (Fig. 2, Ref. 2)
     ISSN: 0287-3796
CY
     Japan
DT
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ΙΑ
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     ANSWER 125 OF 154 LIFESCI
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TI
     Linkage and physical mapping of the porcine prepro- ***orexin***
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     Malek, M.; Marklund, S.; Dyer, C.; Matteri, R.; Rothschild, M. Department of Animal Science, 2255H Kildee Hall, Iowa State University,
ΑU
CS
     Ames, IA 50011, USA; E-mail: mfrothsc@iastate.edu
S0
     Mammalian Genome [Mamm. Genome], ( ***20000400*** ) vol. 11, no. 4, pp.
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DT
     Journal
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L5
     ANSWER 126 OF 154
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        ***Orexin*** --a view discovery in obese research.
TI
ΑU
     Sheng li ke xue jin zhan [Progress in physiology], (1) 47-9. Ref: 9
S0
                                                             ***(2000 Jan)***
                                                                                    31
     Journal code: 20730140R. ISSN: 0559-7765.
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           ***hypocretins*** / ***orexins*** : novel hypothalamic
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de Lecea L; Sutcliffe J G
      Department of Molecular Biology, The Scripps Research Institute, La Jolla, California 92037, USA.. llecea@scripps.edu
CS
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      Cellular and molecular life sciences: CMLS,
                                                           ***(1999 Oct 30)***
      (5-6) 473-80.
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      Journal code: 9705402. ISSN: 1420-682x.
CY
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TI
      Forty winks: molecular basis of sleep disorders.
AU
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     Molecular medicine today,
                                     ***(2000 Dec)***
SO
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      Journal code: 9508560. ISSN: 1357-4310.
CY
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            ***orexin***
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                             OX1
                                    ***receptor***
                                                        activates a novel Ca2+ influx
     pathway necessary for coupling to phospholipase C.
     Lund P E; Shariatmadari R; Uustare A; Detheux M; Parmentier M; Kukkonen J
ΑU
     P; Akerman K E
CS
     Department of Physiology, Division of Cell Physiology, Uppsala University,
     Biomedical Centre (BMC), P.O. Box 572, S-75123 Uppsala, Sweden.
Journal of biological chemistry, ***(2000 Oct 6)*** 275 (40
S0
                                                                     275 (40) 30806-12.
     Journal code: 2985121R. ISSN: 0021-9258.
CY
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                              MEDLINE on STN
ΑN
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DN
     PubMed ID: 10657511
ΤI
     Two important systems in energy homeostasis: melanocortins and
     melanin-concentrating hormone.
     Tritos N A; Maratos-Flier E
     Joslin Diabetes Center, Boston, MA, 02215, USA. Neuropeptides, ***(1999 Oct)*** 33 (5) 339-
CS
S0
     Neuropeptides, ***(1999 Oct)*** 33
Journal code: 8103156. ISSN: 0143-4179.
                                              33 (5) 339-49. Ref: 98
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        ***Hypocretin***
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CM
      Comment in: Lancet. 2000 Apr 8;355(9211):1274-5. PubMed ID: 10770327
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      Nishino S; Ripley B; Overeem S; Lammers G J; Mignot E
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NC
      HL59601 (NHLBI)
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      NS33797 (NINDS)
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      Journal code: 2985213R. ISSN: 0140-6736.
CY
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         ***Orexins***
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      Orexiny a orexinove receptory.
      Kotaska K; Prusa R
Ustav klinicke biochemie a patobiochemie 2. LF UK, Praha.
"**(1999 Aug)***
ΑU
CS
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      48 (3) 119-21. Řef: 17
Journal code: 2984710R. ISSN: 1210-6313.
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                                    ***hypocretins***
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     Narcolepsy: a key role for
                                                         ( ***orexins***
     Comment on: Cell. 1999 Aug 20;98(4):437-51. PubMed ID: 10481909
CM
     Comment on: Cell. 1999 Aug 6;98(3):365-76. PubMed ID: 10458611
     Siegel J<sub>M</sub>
ΑU
CS
     Neurobiology Research, Veterans Administration Medical Center, North
     Hills, California 91343, USA.
Cell, ***(1999 Aug 20)*** 98 (4) 409-12.
S<sub>0</sub>
     Journal code: 0413066. ISSN: 0092-8674.
CY
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DT
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     Novel neurotransmitters for sleep and energy homeostasis.
ΑU
     Sutcliffe J G; de Lecea L
     Department of Molecular Biology, Scripps Research Institute, La Jolla,
CS
     California 92037, USA.
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MH58543 (NIMH)
NS33396 (NINDS)
      Results and problems in cell differentiation,
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      Journal code: 0173555. ISSN: 0080-1844.
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      1999106186
ΑN
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DN
      PubMed ID: 9889512
         ***Orexins*** --new hypothalamic peptides that stimulate appetite].
TI
      Orexiner--nya hypotalamiska peptider som stimulerar aptit.
ΑU
      Meister B; Hakansson M L
      Institutionen for neurovetenskap, Karolinska institutet, Stockholm. Lakartidningen, ***(1998 Dec 16)*** 95 (51-52) 5885-7. Ref: 10
CS
SO
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      Entered Medline: 19990125
      ANSWER 136 OF 154 PASCAL COPYRIGHT 2004 INIST-CNRS. ALL RIGHTS
L5
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      Copyright .COPYRGT. 2000 INIST-CNRS. All rights reserved.
      Reciprocal relation of food intake and sympathetic activity:
TIEN
      experimental observations and clinical implications
      Endocrinology of obesity: basic, clinical and therapeutic aspects
ΑU
      BRAY G. A.
      PASQUALI Renato (ed.)
      Pennington Biomedical Research Center, Baton Rouge, LA 70808, United
CS
      International journal of obesity. Supplement,
SO.
                                                         ***(2000)***
                                                                         , 24(2),
      S8-S17, 101 refs.
      Conference: Endocrinology of Obesity: Basic, Clinical and Therapeutic
      Aspects. Satellite Symposium, Venice (Italy), Sep 1998
      ISSN: 1359-6373
DT
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BL
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CY
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LA
      INIST-18243S, 354000090663260030
ΑV
L5
      ANSWER 137 OF 154 PHARMAML COPYRIGHT 2004 MARKETLETTER ON STN
AN
      1648228
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ΤI
      New drug discovery: is genomics delivering? asks Lehman Brothers
so
      Marketletter August 16, 1999
DT
      Newsletter
WC
      1420
L5
     ANSWER 138 OF 154 PHIN COPYRIGHT 2004 PJB on STN
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     New research offers hope in sleep
TI
     Scrip ( ***1999*** ) No. 2465 p21
SO
DT
     Newsletter
FS
     FULL
L5
     ANSWER 139 OF 154 PROMT COPYRIGHT 2004 Gale Group on STN
```

ACCESSION NUMBER: 2000:889662 PROMT TITLE: SCIÈNCE SCAN RECOMBINANT DRUG SAVED LIVES, LIMBS OF SEPTIC SHOCK PATIENTS IN LARGE CONTROLLED CLINICAL TRIAL. SOURCE: BIOWORLD Today, (***16 Oct 2000***) Vol. 11, No. 200. PUBLISHER: American Health Consultants, Inc. DOCUMENT TYPE: Newsletter LANGUAGE: English WORD COUNT: 958 *FULL TEXT IS AVAILABLE IN THE ALL FORMAT* L5 ANSWER 140 OF 154 PROMT COPYRIGHT 2004 Gale Group on STN ACCESSION NUMBER: 2000:773694 PROMT TITLE: Neurocrine Biosciences, Inc. Announces Nature Medicine Narcolepsy Publication; Absence of ***Hypocretin*** is. Seen to be a Cause of Narcolepsy. Business Wire, (***29 Aug 2000*** SOURCE:) pp. 1023. **PUBLISHER:** Business Wire Newsletter DOCUMENT TYPE: LANGUAGE: English WORD COUNT: 855 *FULL TEXT IS AVAILABLE IN THE ALL FORMAT* ANSWER 141 OF 154 PROMT COPYRIGHT 2004 Gale Group on STN ACCESSION NUMBER: 2000:642390 PROMT Study of Fat-Reducing Protein Opens New Path Toward Obesity TITLE: Treatment. PR Newswire, (***26 Jul 2000***) pp. 8389. SOURCE: **PUBLISHER:** PR Newswire Association, Inc. DOCUMENT TYPE: Newsletter LANGUAGE: English WORD COUNT: 759 *FULL TEXT IS AVAILABLE IN THE ALL FORMAT* L5 ANSWER 142 OF 154 PROMT COPYRIGHT 2004 Gale Group on STN 1999:595824 PROMT ACCESSION NUMBER: SmithKline Beecham Scientists Identify TITLE: ***Receptor*** For Potent Vasoconstricting Hormone. SOURCE: PR Newswire, (***15 Sep 1999***) pp. 8053. **PUBLISHER:** PR Newswire Association, Inc. DOCUMENT TYPE: Newsletter LANGUAGE: English WORD COUNT: 657 *FULL TEXT IS AVAILABLE IN THE ALL FORMAT* L5 ANSWER 143 OF 154 PROMT COPYRIGHT 2004 Gale Group on STN ACCESSION NUMBER: 1999:567548 PROMT TITLE: Narcolepsy Gene Identified in Dogs. Applied Genetics News, (***August 1999***) Vol. 20, No. SOURCE: 1. ISSN: 0271-7107.

PUBLISHER: Business Communications Company, Inc.

DOCUMENT TYPE: Newsletter LANGUAGE: English

WORD COUNT: 273

FULL TEXT IS AVAILABLE IN THE ALL FORMAT

L5 ANSWER 144 OF 154 PROMT COPYRIGHT 2004 Gale Group on STN

ACCESSION NUMBER: 1999:547980 PROMT

TITLE: New drug discovery: is genomics delivering? asks Lehman

Brothers.

SOURCE: Marketletter, Marketletter, (ISSN: 0951-3175 ***23 Aug 1999***) .

PUBLISHER: Marketletter Publications Ltd.

DOCUMENT TYPE: Newsletter

English LANGUAGE: WORD COUNT: 1417

FULL TEXT IS AVAILABLE IN THE ALL FORMAT

ANSWER 145 OF 154 PROMT COPYRIGHT 2004 Gale Group on STN L5

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ACCESSION NUMBER:
                     1999:511958 PROMT
TITLE:
                     TOO MUCH SLEEP CAN BE HAZARDOUS TO HEALTH.
AUTHOR(S):
                     Leff, David N.
                     BIOWORLD Today, ( ***10 Aug 1999*** ) vol. 10, No. 153.
SOURCE:
PUBLISHER:
                     American Health Consultants, Inc.
DOCUMENT TYPE:
                     Newsletter
LANGUAGE:
                     English
WORD COUNT:
                     1006
                     *FULL TEXT IS AVAILABLE IN THE ALL FORMAT*
L5
     ANSWER 146 OF 154 PROMT COPYRIGHT 2004 Gale Group on STN
ACCESSION NUMBER:
                     1999:500436 PROMT
TITLE:
                     UT Southwestern Researchers Create Mice With Narcolepsy.
SOURCE:
                     PR Newswire, ( ***5 Aug 1999*** ) pp. 6759.
PUBLISHER:
                     PR Newswire Association, Inc.
DOCUMENT TYPE:
                     Newsletter
LANGUAGE:
                     English
WORD COUNT:
                     385
                     *FULL TEXT IS AVAILABLE IN THE ALL FORMAT*
     ANSWER 147 OF 154 PROMT COPYRIGHT 2004 Gale Group on STN
L5
ACCESSION NUMBER:
                     1999:497515 PROMT
TITLE:
                     Stanford Researchers Nab Gene for Sleep Disorder.
SOURCE:
                     Business Wire, ( ***5 Aug 1999*** ) pp. 312.
PUBLISHER:
                     Business Wire
DOCUMENT TYPE:
                     Newsletter
                     Enalish
LANGUAGE:
WORD COUNT:
                     1146
                     *FULL TEXT IS AVAILABLE IN THE ALL FORMAT*
     ANSWER 148 OF 154 SCISEARCH COPYRIGHT (c) 2004 The Thomson Corporation.
L5
ΑN
     2000:104862 SCISEARCH
GA
     The Genuine Article (R) Number: 279MU
       ***Hypocretin***
TT
                              ***orexin*** ) deficiency in
                                                               ***human***
     narcolepsy
AU
     Nishino S; Ripley B; Overeem S; Lammers G J; Mignot E (Reprint)
     STANFORD UNIV, SCH MED, DEPT PSYCHIAT, CTR NARCOLEPSY, STANFORD, CA 94305
CS
     (Reprint); STANFORD UNIV, SCH MED, DEPT PSYCHIAT, CTR NARCOLEPSY,
     STANFORD, CA 94305; LEIDEN UNIV, MED CTR, DEPT NEUROL & CLIN NEUROPHYSIOL,
     NL-2300 RC LEIDEN, NETHERLANDS
CYA
     USA; NETHERLANDS
     LANCET, ( ***1 JAN 2000*** ) Vol. 355, No. 9197, pp. 39-40. Publisher: LANCET LTD, 84 THEOBALDS RD, LONDON WC1X 8RR, ENGLAND.
SO
     ISSN: 0140-6736
DT
     Article; Journal
     LIFE; CLIN
FS
LA
     English
REC
     Reference Count: 5
     *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS*
L5
     ANSWER 149 OF 154 SCISEARCH COPYRIGHT (c) 2004 The Thomson Corporation.
AN
     1999:853265 SCISEARCH
GA
     The Genuine Article (R) Number: 252ER
     Stimulation of feeding behavior and food consumption in the goldfish,
     Carassius auratus, by
                              ***orexin*** -A and
                                                      ***orexin***
     Volkoff H; Bjorklund J M; Peter R E (Reprint)
ΑU
     UNIV ALBERTA, DEPT BIOL SCI, EDMONTON, AB T6G 2E9, CANADA (Reprint); UNIV
CS
     ALBERTA, DEPT BIOL SCI, EDMONTON, AB T6G 2E9, CANADA
CYA
                        ***6 NOV 1999*** ) Vol. 846, No. 2, pp. 204-209.
     BRAIN RESEARCH, (
     Publisher: ELSEVIER SCIENCE BV, PO BOX 211, 1000 AE AMSTERDAM,
     NETHERLANDS.
     ISSN: 0006-8993
DT
     Article; Journal
FS
     LIFE
LA
     English
REC
     Reference Count: 25
     *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS*
L5
     ANSWER 150 OF 154 USPATFULL ON STN
       2003:279117 USPATFULL
ΑN
TI
       Hypothalamus-specific polypeptides
```

```
IN
        Sutcliffe, J. Gregor, Cardiff, CA, United States
        Gautvik, Kaare M., Oslo, NORWAY
        De Lecea, Luis, Del Mar, CA, United States
        Bloom, Floyd E., San Diego, CA, United States
        Danielson, Patria E., San Diego, CA, United States
       Gautvik, Vigdis T., Oslo, NORWAY
Kilduff, Thomas S., Menlo Park, CA, United States
Foye, Pamela E., San Diego, CA, United States
PA
        The Scripps Research Institute, La Jolla, CA, United States (U.S.
        corporation)
PT
        US 6635479
                             в1
                                  20031021
        wo 9805352
                     19980212
                                                                           <--
        US 1999-230896
AΤ
                                  19990202 (9)
        wo 1997-us13657
                                  19970801
        US 1996-23220P
PRAI
                              19960802 (60)
        Utility
DT
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LN.CNT
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INCL
        INCLM: 435/325.000
        INCLS: 435/320.100; 536/023.100; 536/023.500
NCL
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        NCLS:
               435/320.100; 536/023.100; 536/023.500
IC
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        ICS: C12N015-00; C12N015-09; C12N005-00; C12N005-02 536/23.1; 536/22.1; 536/23.5; 530/300; 530/350; 435/320.1; 435/325
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 151 OF 154 USPATFULL ON STN
L5
ΑN
        2003:13320 USPATFULL
                 ***orexin***
TI
                                    ***receptor***
                                                       antagonists
        Irving, Elaine Alison, Bengeo, UNITED KINGDOM
IN
        Sanger, Gareth John, Sawbridgeworth, UNITED KINGDOM
        SmithKline Beecham p.l.c., Brentford, UNITED KINGDOM (non-U.S.
PA
        corporation)
PI
        US 6506774
                            в1
                                  20030114
       WO 2000047284 20000817
ΔΤ
        US 2001-913230
                                  20011130 (9)
       WO 2000-EP1147
                                  20000210
                             19990212
PRAI
        GB 1999-3265
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DT
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LN.CNT 405
INCL
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        [7]
        ICM: A61K031-47
EXF
        514/311; 514/312; 514/313
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L5
     ANSWER 152 OF 154 WPIDS COPYRIGHT 2004 THE THOMSON CORP ON STN
     2000-588952 [56]
AN
                          WPIDS
DNC
     C2000-175936
     Obesity treatment comprises administering N-(4-phenoxyphenyl)-oxamic acid
     derivatives or related compounds.
DC
     B05
IN
     CORNELIUS, P; HARGROVE, D M; MORGAN, B P; SWICK, A G; HARGROVE, D
PA
      (PFIZ) PFIZER PROD INC; (CORN-I) CORNELIÚS P: (HARG-Í) HARGROVÉ D:
      (MORG-I) MORGAN B P; (SWIC-I) SWICK A G; (PFIZ) PFIZER INC
CYC
     33
PΙ
     EP 1036564
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                                                    22
                                                           A61K031-225
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A61K031-24

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B1 20020205 (200211)
A1 20020321 (200224)#
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       US 2002035153
                                                                A61K031-195
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                         B2 20030429 (200331)
B 20030124 (200339)
                                                                 A61K031-24
       KR 368354
                                                                 A61K031-196
       CA 2299972
                             20030819 (200357)
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      EP 1036564 A1 EP 2000-300830 20000203; AU 2000016353 A AU 2000-16353
 ADT
       20000211; CA 2299972 A1 CA 2000-2299972 20000229; JP 2000256190 A JP
       2000-49507 20000225; HU 2000000921 A2 HU 2000-921 20000228; KR 2001006712
      A KR 2000-9860 20000228; ZA 2000001000 A ZA 2000-1000 20000229; NZ 503122 A NZ 2000-503122 20000229; US 6344481 B1 Provisional US 1999-122015P 19990301, US 2000-488110 20000120; US 2002035153 A1 Div ex US 2000-488110 20000120, US 2001-978980 20011016; US 6555578 B2 Provisional US 1999-122015P 19990301 Div ex US 2000-488110 20000120 US 2001-978980 20011016; US 6555578 B2 Provisional US 1999-122015P 19990301 Div ex US 2000-488110 20000120 US 2001-978980
       1999-122015P 19990301, Div ex US 2000-488110 20000120, US 2001-978980
       20011016; KR 368354 B KR 2000-9860 20000228; CA 2299972 C CA 2000-2299972
       20000229
      US 6555578 B2 Div ex US 6344481; KR 368354 B Previous Publ. KR 2001006712
 FDT
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            A61K031-192; A61K031-235; A61K045-00; A61P003-04; A61P043-00;
            C07D000-00
 15
       ANSWER 153 OF 154 WPIDS
                                    COPYRIGHT 2004 THE THOMSON CORP ON STN
      2000-105551 [09]
 AN
                             WPIDS
DNC
      C2000-031609
      New phenyl urea and phenylthio urea derivatives useful as
                                                                             ***orexin***
      antagonists for treating e.g. obesity, insomnia, schizophrenia, manic
       depression and diabetes.
DC
       R02
      JOHNS, A; PORTER, R A
ΙN
       (SMIK) SMITHKLINE BEECHAM PLC
PA
CYC
      87
PT
      wo 9958533
                         A1 19991118 (200009)* EN
                                                         33
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          RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW NL
               OA PT SD SE SL SZ UG ZW
           W: AE AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES FI GB
               GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ LC LK LR LS LT LU
               LV MD MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR
              TT UA UG US UZ VN YU ZA ZW
      AU 9940377
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                         A1 20010214 (200111)
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                                                                A61K031-435
      EP 1075478
                         B1 20030416 (200328)
                                                   FN
                                                                C07D471-04
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                         E 20030522 (200341)
      DE 69906960
                                                                C07D471-04
      JP 2003522101
                            20030722 (200350)
                         W
                                                                C07D471-04
      ES 2196806
                        T3 20031216 (200413)
                                                                C07D471-04
      WO 9958533 A1 WO 1999-EP3100 19990504; AU 9940377 A AU 1999-40377
      19990504; EP 1075478 A1 EP 1999-923540 19990504, WO 1999-EP3100 19990504
      US 6372757 B1 WO 1999-EP3100 19990504, US 2000-700002 20001108; EP 1075478
      B1 EP 1999-923540 19990504, WO 1999-EP3100 19990504; DE 69906960 E DE 1999-606960 19990504, EP 1999-923540 19990504, WO 1999-EP3100 19990504;
      2003522101 w wo 1999-EP3100 19990504, JP 2000-548337 19990504; ES 2196806
      T3 EP 1999-923540 19990504
      AU 9940377 A Based on WO 9958533; EP 1075478 A1 Based on WO 9958533; US
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      6372757 B1 Based on WO 9958533; ÉP 1075478 B1 Based on WO 9958533; DE
      69906960 E Based on EP 1075478, Based on WO 9958533; JP 2003522101 W Based
      on WO 9958533; ES 2196806 T3 Based on EP 1075478
PRAI GB 1999-3268
                               19990212; GB 1998-9972
                                                                   19980508;
      GB 1998-9988
                               19980508
           A61K031-435; C07D471-04
A61K031-4375; A61K031-44; A61P003-04; A61P015-00; A61P025-00;
IC
      ICM
      ICS
            A61P025-04; A61P025-18; A61P025-20; A61P025-22; A61P025-24;
            A61P043-00
     C07D221:00; C07D221:00, C07D471-04; C07D221:00, C07D471-04; C07D221:00,
ICI
            C07D471-04
L5
      ANSWER 154 OF 154 WPIDS
                                   COPYRIGHT 2004 THE THOMSON CORP ON STN
      1999-315250 [27]
ΑN
                            WPIDS
DNC
      C1999-093223
      Composition for treating obesity and diabetes comprises a specific beta-3
TI
      agonist and an anorectic agent.
DC
      B02 B03 C02
```

us 6344481

DOW, R L

IN

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PA
      (PFIZ) PFIZER PROD INC
CYC
      30
ΡI
      EP 920864
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A 19990824 (199944)
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      KR 99062718
                          A 19990726 (200043)
                                                                    A61K031-44
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ADT
      EP 920864 A1 EP 1998-309273 19981112; AU 9896055 A AU 1998-96055 19981202;
      HU 9802795 A2 HU 1998-2795 19981202; JP 11228447 A JP 1998-335819 19981126; CA 2255318 A1 CA 1998-2255318 19981201; KR 99062718 A KR 1998-52532 19981202
PRAI US 1997-67268P
                                19971203
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ICS A61K031-00; A61K031-13; A61K038-00; A61K038-22
STN INTERNATIONAL LOGOFF AT 15:56:22 ON 18 OCT 2004
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